Effects of $I_{Kur}$ blocker MK-0448 on human right atrial action potentials from patients in sinus rhythm and in permanent atrial fibrillation

Simone Loose1, Judith Mueller1, Erich Wettwer1, Michael Knaut2, John Ford3, James Milnes3 and Ursula Ravens1*

1 Department of Pharmacology and Toxicology, Medical Faculty Carl Gustav Carus, Dresden University of Technology, Dresden, Germany
2 Clinic for Cardiac Surgery, Heart Center Dresden, Dresden, Germany
3 Xention Ltd., Cambridge, UK

INTRODUCTION

In search of safer antiarrhythmic drugs for the treatment of atrial fibrillation (AF), selective inhibitors of $I_{Kur}$ have been designed as atrial-selective class III antiarrhythmic agents with therapeutic potential in AF. Since they target the Kv1.5 channels which are functional only in atria (Mays et al., 1995), these agents are considered to be devoid of proarrhythmic effects in ventricles and hence have received a lot of interest in the scientific community (for review see Ford and Milnes, 2008; Tamargo et al., 2009; Ravens, 2010; Ravens and Wettwer, 2011).

We have previously investigated the electrophysiological effects of several $I_{Kur}$ inhibitors, including 4-aminopyridine, AVE0118, XEN-D0101, all of which elevated the plateau phase of human right atrial action potentials (APs) recorded in trabeculae of atrial appendage obtained from patients in sinus rhythm (SR) or AF (Wettwer et al., 2004; Christ et al., 2008; Ford et al., 2013). Interestingly, action potential duration (APD90) and effective refractory period (ERP) were shortened in SR but prolonged in AF preparations.

Recently, first-time-in-man results with another selective Kv1.5 inhibitor, MK-0448 were published (Pavri et al., 2012). MK-0448 failed to affect relative atrial ERP in healthy volunteers, and the authors concluded that $I_{Kur}$ block cannot be expected to exert therapeutically useful antiarrhythmic effects in AF. Since nothing is known about the effects of MK-0448 in ex-vivo human atrial tissue, we investigated its effects on the shapes of atrial APs and ERP with standard microelectrode techniques in trabeculae isolated from right atrial appendages of patients in SR and AF, who had to undergo open heart surgery. In SR preparations, MK-0448 elevated the plateau potential and shortened APD and ERP at 1 Hz as expected from $I_{Kur}$ blockers. We also confirm that MK-0448 effectively blocks Kv1.5 current in mouse fibroblasts stably expressing the hKCNA5 gene.

METHODS

MOUSE FIBROBLASTS

HK2BN9 cells expressing human Kv1.5 channels were kindly provided by Dr. Tamkun (Snyders et al., 1993). The cells were cultured in Dulbecco's modified eagle medium containing 10% fetal calf serum and 1% penicillin/streptomycin under atmospheric conditions of 5% CO2 at 37°C. 250 μg/ml G418 was added to the medium for selection. Expression of Kv1.5 was induced by treating the cells with 1 μM dexamethasone 24 h before an experiment. The cells were studied with a standard single electrode whole-cell voltage-clamp set-up as described previously (Radicke et al., 2006). The bath solution contained (in mM): NaCl 150, KCl 5.4, MgCl2 2, CaCl2 2.0, HEPES 10, glucose 10, pH 7.4, temperature 25°C. The electrode solution contained (in mM): KCl 40, potassium aspartate 80, NaCl 8, CaCl2 2, MgATP 5, EGTA 5, GTP 0.1, HEPES 10, pH 7.4 adjusted with KOH. Series resistances were between 5 and 10 MΩ and were compensated by up to 85%. Cell capacitance was measured with small hyperpolarizing clamp steps from −40 mV to −42 mV, average value was 20.8 ± 1.2 pA/pF ($n = 33$) and did not differ between cells tested at 0.5 and 3 Hz. Voltage clamp pulse generation, data collection and analysis were performed with the ISO2 software (MFK, Niedernhausen, Germany). Data were not corrected for the junction potential.

Keywords: MK-0448, atrial fibrillation, Kv1.5, $I_{Kur}$ blocker, human right atrial action potentials
which was 11.7 mV, as calculated with the JPCalc software (P.H. Barry, Sydney, Australia).

PATIENTS’ DIAGNOSES AND MEDICATIONS

Atrial tissue was obtained from patients receiving cardiac surgery because of coronary artery or valve disease. All patients gave written informed consent. The study was approved by the ethic committee of the Medical Faculty of Dresden University of Technology (No. EK790799). The patients are listed in Table 1. The study included 13 drug-exposed and 9 TMC preparations from patients in sinus rhythm (SR group), 9 drug-exposed and 9 TMC preparations from patients with permanent atrial fibrillation (AF group, defined as permanent atrial fibrillation for ≥6 months at time of tissue collection), and 3 drug-exposed preparations from patients with intermittent AF (episodes of spontaneously terminating AF lasting for a few seconds up to several hours). The medication of the patients typically included ACE-inhibitors, β-blockers, nitrates, lipid-lowering drugs, and diuretics.

EXPERIMENTAL SET-UP

Small pieces (50–100 mg in weight) of human right atrial appendages were transported to the laboratory in a special Ca²⁺-free transport solution at 20–25°C, composition in mM: 100 NaCl, 10 KCl, 1.2 KH₂PO₄, 5 MgSO₄, 50 taurine, 5 MOPS, 30 2,3-butanediol monoxime (BDM), pH 7.0. Either free-running trabeculae (<1 mm in diameter and 2–5 mm in length) or trabeculae together with attached atrial wall were dissected and mounted on the bottom of a 5 ml organ bath perfused with 50 ml of recirculating, oxygenized Tyrode’s solution at a flow rate of 7 ml/min at 36 ± 1°C (composition in mM: 126.7 NaCl, 0.42 NaH₂PO₄, 22 NaHCO₃, 5.4 KCl, 1.8 CaCl₂, 1.5 MgCl₂, pH 7.4 when equilibrated with 5% CO₂ in O₂). Preparations were electrically stimulated at a basal rate of 1 Hz with isolated square-wave stimuli of 1 ms duration, 2 times threshold intensity. Transmembrane potentials were recorded with glass microelectrodes filled with 2.5 M KCl. Tip resistances of the electrodes were between 20 and 80 MΩ. Both timing of the driving stimuli and preprocessing of the transmembrane potential responses were carried out with a computer-aided action potential recording system (Wettwer et al., 2004). Chart for Windows, Version 7.3.5, recording programme (ADInstruments Pty Ltd., Castle Hill, Australia) was used for analysis of action potential parameters.

Each experiment was preceded by a 60 min equilibration period during which the preparations were allowed to stabilize and residual BDM was completely washed out. Numerical values of the main action potential parameters had to remain constant, i.e., within a range of ±5% for at least 10 min. When the microelectrode moved out of the cell, a new impalement was made at the same place and the experiment was continued if the new recording did not differ from the old one by visual judgment and showed less than 3% difference upon final numerical analysis.

MEASUREMENT OF EFFECTIVE REFRACTORY PERIOD

Effective refractory period (ERP) was measured by a single extra stimulus interpolated after trains of 10 regular stimulation pulses at the basal frequency of 1 Hz with decreasing intervals in steps of 5 ms from the last regular pulse until the extra stimulus failed to elicit an action potential. Following a short recovery period the preparation was exposed to MK-0448 (3 μM) at 1 Hz for another 60 min and ERP was tested once more in the presence of drug.

STOCK SOLUTIONS AND DRUG APPLICATION

MK-0448 was synthesized by Xention Ltd. Cambridge UK. In a few experiments, MK-0448 was applied in cumulatively increasing concentrations by directly adding aliquots from a 10 mM stock solution with DMSO as solvent to 50 ml of the circulating bath medium. The DMSO concentration never exceeded 0.2%, a concentration which did not induce any significant change when applied alone. Preparations were exposed to each concentration which did not induce any significant change when applied alone. Preparations were exposed to each concentration of MK-0448 for 20 min.

ANALYSIS OF ACTION POTENTIAL PARAMETERS

The following standard parameters of atrial APs were analyzed: resting membrane potential, RMP (mV), action potential amplitude, APA (mV), action potential duration at 20, 50, and 90%
of repolarization, APD$_{20}$, APD$_{50}$ and APD$_{90}$ (ms), maximum rate of depolarization, $dV/dt_{\text{max}}$ (V/s) and the “plateau potential” defined as the mean absolute membrane potential (mV) in the time window between 20 and 30% of APD$_{90}$ (PLT$_{20}$). Preparations with resting potentials less than $-65\,\text{mV}$ and amplitudes below 80 mV were discarded.

**STATISTICAL ANALYSIS**

GraphPad Prism (version 4.03) was used for statistical analysis. In the concentration-response curves (see Table 2) the values obtained at increasing concentrations were compared to pre-drug controls by One-Way ANOVA followed by Bonferroni’s multiple comparison test. The individual action potential parameters of SR and permanent AF preparations in the presence of 3 $\mu$M MK-0448 were compared to pre-drug controls by paired Student’s $t$-test (see Table 3). Differences are considered statistically significant for $P < 0.05$.

**RESULTS**

**EFFECTS OF MK-0448 ON Kv1.5 CURRENT**

Kv1.5 currents, which was measured during step potential changes from a holding potential of $-60\,\text{mV}$ to various test potentials between $-40$ and $+80\,\text{mV}$ had a threshold potential of $-30\,\text{mV}$, activated rapidly and hardly inactivated within 500 ms (data not shown). For drug testing we chose 50 ms duration voltage-clamp steps to $+50\,\text{mV}$ that elicited mean current amplitudes of 85.0 ± 10.5 pA/pF at 0.5 Hz ($n = 16$) and 110.6 ± 16.9 pA/pF at 3 Hz $n = 17$, $P = 0.2133$. MK-0448 (100 nM) blocked Kv1.5 current amplitude in a frequency-dependent manner (Figure 1A). At 0.5 Hz, the block reached a steady-state within 5–10 min of drug application and partially recovered during a 5-min period during which the cell was continuously clamped at $-60\,\text{mV}$ (Figure 1B). When cells were exposed to MK-0448 during a 5 min period held at $-60\,\text{mV}$, to maintain channels in the closed-state, $I_{\text{Kv1.5}}$ of the first post-rest pulse was not different from pre-rest. Subsequent pulses in the

| Table 2 | Concentration-dependent effects of MK-0448 at 1, 3, and 10 $\mu$M (1 Hz). |
|-----------------|-----------------|-----------------|-----------------|-----------------|
| **SR** | **Control** | **1.0 $\mu$M** | **3.0 $\mu$M** | **10.0 $\mu$M** |
| **APD$_{90}$ (ms)** | $307.6 \pm 10.2$ | $269.7 \pm 8.2$ | $266.9 \pm 12.2$ | $260.9 \pm 19.5$ | $271.6 \pm 13.7$ |
| **APD$_{50}$ (ms)** | $116.3 \pm 28.6$ | $136.5 \pm 33.1$ | $149.6 \pm 35.6$ | $167.4 \pm 20.9$* | $154.6 \pm 17.4$ |
| **APD$_{20}$ (ms)** | $5.4 \pm 1.5$ | $51.3 \pm 20.5$ | $92.0 \pm 23.2$ | $92.5 \pm 23.8$** | $63.7 \pm 25.3$ |
| **PLT$_{20}$ (mV)** | $-20.4 \pm 3.4$ | $-4.0 \pm 6.2$*** | $+2.8 \pm 6.9$*** | $+3.4 \pm 6.8$*** | $-6.0 \pm 6.7$ |
| **APA (mV)** | $95.3 \pm 2.7$ | $94.5 \pm 2.3$ | $93.2 \pm 3.8$ | $88.6 \pm 4.7$ | $86.5 \pm 7.2$ |
| **RMP (mV)** | $-74.2 \pm 1.9$ | $-74.5 \pm 4.0$ | $-72.8 \pm 3.5$ | $-70.9 \pm 4.6$ | $-70.0 \pm 5.3$ |
| **dV/dt$_{\text{max}}$ (V/s)** | $300.8 \pm 24.4$ | $267.3 \pm 29.8$ | $269.2 \pm 27.9$ | $191.8 \pm 58.3$ | $186.3 \pm 54.3$ |

| **Permanent AF** | **Control** | **1.0 $\mu$M** | **3.0 $\mu$M** | **10.0 $\mu$M** |
|-----------------|-----------------|-----------------|-----------------|-----------------|
| **APD$_{90}$ (ms)** | $190.8 \pm 11.8$ | $200.7 \pm 9.3$ | $215.2 \pm 11.1$ | $221.4 \pm 15.2$* | $222.9 \pm 17.0$ |
| **APD$_{50}$ (ms)** | $101.6 \pm 7.4$ | $125.7 \pm 7.5$ | $143.1 \pm 8.4$* | $147.8 \pm 11.1$** | $145.9 \pm 11.8$ |
| **APD$_{20}$ (ms)** | $31.8 \pm 8.8$ | $55.7 \pm 4.0$ | $73.4 \pm 5.8$** | $78.2 \pm 7.1$** | $71.7 \pm 8.4$ |
| **PLT$_{20}$ (mV)** | $-9.9 \pm 2.4$ | $3.0 \pm 1.8$* | $9.5 \pm 2.2$*** | $11.6 \pm 4.3$*** | $4.7 \pm 4.2$ |
| **APA (mV)** | $100.9 \pm 3.6$ | $104.0 \pm 2.2$ | $101.4 \pm 3.7$ | $98.2 \pm 6.7$ | $96.9 \pm 5.4$ |
| **RMP (mV)** | $-79.2 \pm 3.3$ | $-78.9 \pm 2.6$ | $-75.7 \pm 3.0$ | $-70.9 \pm 3.9$*** | $-74.6 \pm 2.4$ |
| **dV/dt$_{\text{max}}$ (V/s)** | $211.0 \pm 36.4$ | $204.0 \pm 40.9$ | $177.8 \pm 23.6$ | $147.3 \pm 32.2$ | $200.1 \pm 37.2$ |

MK-0448, APD$_{90}$, APD$_{50}$, APD$_{20}$, action potential duration at 90, 50, and 20% of repolarization, respectively; PLT$_{20}$, average plateau potential at 20–30% of APD$_{90}$; APA, action potential amplitude; RMP, resting membrane potential; dV/dt$_{\text{max}}$, maximum rate of depolarization. Means ± s.e.m. from 5 experiments in which all concentrations indicated were tested.

$^{*}P < 0.05$, $^{**}P < 0.01$, $^{***}P < 0.001$; One-Way-ANOVA followed by Bonferroni’s multiple comparison test (drug vs. pre-drug control).
FIGURE 1 | Effects of MK-0448 on Kv1.5 currents measured in HK2BN9 cells at 25°C. (A) Voltage clamp and current traces obtained before (black) and after superfusion of cells with 100 nM MK-0448 (green) for 10 min. Rate of clamp pulses as indicated. (B) Onset of blocking effect of MK-0448 (100 nM) on peak Kv1.5 current and recovery from block during 5 min at holding potential of −60 mV. Individual current traces at top were obtained at the time points indicated by small letters in the diary plot at the bottom. (C) Onset of drug effect following exposure during 5 min of continuous clamping at −60 mV. Step frequency: 3 Hz.

EFFECTS OF MK-0448 ON ACTION POTENTIALS

Action potentials recorded in right atrial trabeculae from patients in SR had the typical spike-and-dome configuration (Figure 2), in permanent AF tissue the shape was triangular and is the result of electrical remodeling [for review see (Dobrev and Ravens, 2003)]. The most striking concentration-dependent effect of MK-0448 (1–10 μM) on SR action potentials at the basal stimulation frequency of 1 Hz was a robust elevation in plateau potential (PLT20) (Figure 2; Table 2). In this set of experiments, the APD90 tended to be shortened (not significant) whereas APD20 and APD50 were prolonged, reaching the level of statistical significance only at the highest concentration (see Table 2). Resting membrane potential (RMP), action potential amplitude (APA) and maximum depolarization velocity (dV/dt max) were not significantly affected by MK-0448 (see Table 2). Before and after drug treatment all preparations followed the stimulation, even at the highest concentration.

The effects of MK-0448 on ERP were studied only at 3 μM. This concentration produced maximum changes within 40–50 min of drug application, and these were partially reversible after 60 min of washout. In 6 preparations from patients in SR, APD90, and ERP were significantly shortened (Figure 3; Table 3). Similar effects were observed in 2 preparations from patients in intermittent AF (Figure 3).

In permanent AF trabeculae, MK-0448 also significantly elevated the plateau potential to a similar extent as in SR, but in contrast to SR, prolonged APD at the three analyzed repolarization levels (Figures 2, 3; Table 3). In permanent AF preparations, APA, RMP, and dV/dt max were also not significantly affected, with exception of RMP with 10 μM MK-0448.

Because of limited available we could only study the effects of MK-0448 in 2 trabeculae from patients in intermittent AF, a typical recording is shown in Figure 3; 3 others were used as TMCs. Similar to the results in SR preparations, MK-0448 shortened APD90 and ERP in the 2 intermittent AF preparations.

In time-matched control experiments with tissue from patients in SR, intermittent AF, and AF, action potential parameters did not change significantly (data not shown).

DISCUSSION

The characteristic changes in action potential parameters observed with MK-0448 in human right atrial trabeculae are consistent with block of Kv1.5 channels as reported previously for several drugs [e.g., 4-AP, (Wang et al., 1993; Amos et al., 1996); DPO-1, (Regan et al., 2006); AVE-0118, (Christ et al., 2008); XEN-D0101, (Ford et al., 2013)].

MK-0448 was described as highly selective for expressed Kv1.5 channels [IC50 ~10 nM; (Pavri et al., 2012)] over a wide variety of standard cardiac ion channels. hERG channels are about 10,000-fold, inward rectifier current Kir2.1 and Kir3.1/3.4 about...
1000-fold, and Kv7.1 (IKs) some 10- to 100-fold less sensitive than Kv1.5 (Pavri et al., 2012). The voltage-clamp data in a heterologous expression system confirm that MK-0448 (100 nM) produced a robust block of human Kv1.5 with a rather slow onset of action and partial recovery from block at −60 mV. Although our results do not allow precise characterization of drug-channel interaction they clearly indicate that resting channels recover from block and activation is necessary for development of block, suggesting preferential binding to open and/or inactivated channels.

In previous experiments, we demonstrated that low concentrations of 4-aminopyridine selectively inhibit \( I_{\text{Kur}} \) in human atrial myocytes (Wettwer et al., 2004). This blocker elevated the plateau potential in SR preparations (EC\(_{50} \sim 15 \mu M\)). In addition,
4-AP at concentrations that are considered to be selective for I_{Kur} block significantly shortened APD_{90} in SR but prolonged APD_{90} in permanent AF trabeculae (Wettwer et al., 2004). The most prominent effects of MK-0448 on human atrial APs, i.e., elevation of plateau potential, shortening of APD_{90} in SR and prolongation of APD_{20} APD_{50} and APD_{90} in permanent AF, are in line with our previous findings with other I_{Kur} inhibitors, so that we are confident to conclude that MK-0448 does indeed block I_{Kur} in human atrial trabeculae.

Unlike inhibition of hERG channels, inhibition of I_{Ks} was not associated with a proarrhythmic ventricular effect in animal models (Lynch et al., 1999). Nevertheless, modulation of I_{Ks} by MK-0448 could also reduce repolarization reserve in the ventricle and become arrhythmogenic in ventricles under appropriate conditions (Lynch et al., 1999). It is important to note that it is difficult to assess the ventricular proarrhythmic risk of a drug that modulates I_{Ks} in healthy volunteers with normal ventricular repolarization reserve. Therefore, the atrial selectivity of MK-0448 would need to be carefully assessed in patients.

The shortening of APD_{90} and ERP by Kv1.5 blockers in SR is proposed to be caused by the marked elevation of the plateau potential in to a voltage range where more hERG channels are activated, and, though inactivated rapidly, can recover from inactivation at more negative potentials and thus hasten final repolarization (Gintant, 2000).

Besides distinct alterations in shapes of action potentials, loss of rate-adaptation of ERP and APD_{90} to frequency has been recognized as a hallmark of atrial remodeling both in patients (e.g., Attuel et al., 1982; Daoud et al., 1996) and in ex vivo studies (Franz et al., 1997; Dobrev and Ravens, 2003). In our present study and in the recent “first-time-in-human” study with MK-0448 all electrophysiological testing was in a frequency range much below that of AF and one could argue that based on our SR data that it is not surprising that MK-0448 did not prolong AERP in healthy volunteers (Pavri et al., 2012).

Experimental and clinical evidence suggests that atrial fibrillation is due to micro-reentry triggered by ectopic foci located in the proximity of the pulmonary veins (Haissaguerre et al., 1982; Daoud et al., 1996). In SR preparations, MK-0448 produced shortening of APD_{90} and ERP but in permanent AF tissue, prominent APD_{90}- and ERP-prolonging effects were observed with the drug. For prevention of persistent AF recurrence, prolongation of ERP by MK-0448 in AF preparations is certainly a highly desirable mode of action that may protect the atrial myocardium during the phase of reverse remodeling after cardioversion although if due to I_{Ks} modulation this may come at a risk of inducing ventricular proarrrhythmia. It remains to be determined if selective I_{Kur} drugs (i) are more effective at prolonging ERP at higher atrial rates, (ii) pharmacologically convert recent onset AF, (iii) reducing AF burden in paroxysmal AF, and (iv) prevent the recurrence of persistent AF.

CONCLUSION

In summary, the I_{Kur} blocker MK-0448 consistently elevates the action potential plateau in human atrial trabeculae irrespective of the initial shapes of the control action potentials and whether the preparations were obtained from patients in SR or permanent AF. In SR preparations, MK-0448 produced shortening of APD_{90} and ERP, but in permanent AF tissue, prominent APD_{90}- and ERP-prolonging effects were observed with the drug. For prevention of persistent AF recurrence, prolongation of ERP by MK-0448 in AF preparations is certainly a highly desirable mode of action that may protect the atrial myocardium during the phase of reverse remodeling after cardioversion although if due to I_{Ks} modulation this may come at a risk of inducing ventricular proarrrhythmia. It remains to be determined if selective I_{Kur} drugs (i) are more effective at prolonging ERP at higher atrial rates, (ii) pharmacologically convert recent onset AF, (iii) reducing AF burden in paroxysmal AF, and (iv) prevent the recurrence of persistent AF.

ACKNOWLEDGMENTS

The authors gratefully acknowledge the financial support of the European Union (FP7-Health T2-2010-261057 “EUTRAF”) and Xention Limited. We wish to thank all patients and the cardiac surgeons Klaus Matschke and Michael Knaut for their invaluable support, and Konstanze Fischer for her expert technical assistance.

REFERENCES

Amos, G. J., Wettwer, E., Metzger, E., Li, Q., Himmel, H. M., and Ravens, U. (1996). Differences between outward currents of human atrial and subepicardial ventricular myocytes. J. Physiol. 491(Pt 1), 31–50.

Attuel, P., Childers, R., Cauchemez, B., Poveda, J., Majrica, I., and Coumel, P. (1982). Failure in the rate adaptation of the atrial refractory period: its relationship to vulnerability. Int. J. Cardiol. 2, 179–197. doi:10.1016/0167-5273(82)90032-8

Christ, T., Wettwer, E., Voigt, N., Hala, O., Radicke, S., Matschke, K., et al. (2008). Pathology-specific effects of the I'(Kur)/(I(to)/(I(K,ACh)) blocker AVE0118 on ion channels in human chronic atrial fibrillation. Br. J. Pharmacol. 154, 1619–1630. doi:10.1038/bjp.2008.209
Haissaguerre, M., Jais, P., Shah, D. C., Takahashi, A., Hocini, M., Quiniou, G., Gintant, G. A. (2000). Characterization and functional consequences of delayed rapid delayed-rectifier current (I_{Kur}): rationale, pharmacology and evidence for potential therapeutic value. J. Cardiovasc. Pharmacol. 36, 105–120. doi: 10.1097/0000385-000043

Ford, J., Milnes, J., Wettwer, E., Christ, T., Rogers, M., Sutton, K., et al. (2013). Ultra-rapid delayed rectifier channels: molecular basis and therapeutic implications. Circ. Res. 89, 776–785. doi: 10.1093/cvr/cvr398

Regan, C. P., Wallace, A. A., Cresswell, H. K., Atkins, C. L., and Lynch, J. J. Jr. (2006). In vivo cardiac electrophysiologic effects of a novel diphenylphosphine oxide IKur blocker, (2-Isopropyl-5-methylcyclohexyl) diphenylphosphine oxide, in rat and nonhuman primate. J. Pharmacol. Exp. Ther. 316, 727–732. doi: 10.1124/jpet.105.094839

Snyders, D. I., Tamkun, M. M., and Bennett, P. B. (1993). A rapidly activating and slowly inactivating potassium channel cloned from human heart. Functional analysis after stable mammalian cell culture expression. J. Gen. Physiol. 101, 513–543. doi: 10.1085/s0014-513

Conflict of Interest Statement: James Milnes and John Ford are employees of Xention Ltd., John Ford is co-founder and holds stock in Xention Ltd., Ursula Ravens has acted as consultant to Xention. Simone Loose, Judith Mueller, Michael Knaut, Erich Wettwer have nothing to disclose.

Received: 27 November 2013; accepted: 12 February 2014; published online: 04 March 2014.

Citation: Loose S, Mueller J, Wettwer E, Knaut M, Ford J, Milnes J and Ravens U (2014) Effects of I_{Kur} blocker MK-0448 on human right atrial action potentials from patients in sinus rhythms and in permanent atrial fibrillation. Front. Pharmacol. 5:26. doi: 10.3389/fphar.2014.00026

This article was submitted to Pharmacology of Ion Channels and Channelopathies, a section of the journal Frontiers in Pharmacology.

Copyright © 2014 Loose, Mueller, Wettwer, Knaut, Ford, Milnes and Ravens. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) or licensor are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.