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Biased Effect of Cardiotonic Steroids on Na/K-ATPase-mediated Signal Transduction

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ABBREVIATIONS:

NKA, Na/K-ATPase; CTS, cardiotonic steroids; Akt, protein kinase B; BSA, bovine serum albumin; DMEM, Dulbecco's Modified Eagle's Medium; ECL, enhanced chemiluminescence; ERK, extracellular-signal related kinase; c-Src, Proto-oncogene tyrosine-protein kinase Src; PI3Ks, Phosphoinositide 3-kinases; mAb, monoclonal antibodies; GPCRs, G protein-coupled receptors; EGFR, epidermal growth factor receptor; PLC- γ , phospholipase C γ ; PKC, protein kinase C; IP3, inositol trisphosphate.

Abstract

Recent studies have revealed that Na/K-ATPase (NKA) can transmit signals through ion pumping-independent activation of pathways relayed by distinct intracellular protein/lipid kinases and endocytosis challenges the traditional definition that cardiotonic steroids (CTS) is NKA inhibitor. While additional effects of CTS have long been suspected, revealing its agonist impact through the NKA receptor could be a novel mechanism in understanding the basic biology of NKA. In this study, we tested whether different structural CTS could trigger different sets of NKA/effector interactions, resulting in biased signaling responses without compromising ion-pumping capacity. Using purified NKA, we found that ouabain, digitoxigenin, and somalin cause comparable levels of NKA inhibition. However, while endogenous ouabain stimulates both protein kinases

and NKA endocytosis, digitoxigenin and somalin bias to protein kinases and endocytosis respectively in LLC-PK1 cells. The positive inotropic effects of CTS are traditionally regarded as NKA inhibitors. However, CTS-induced signaling occurs at concentrations at least one order of magnitude lower than that of inotropy, which eliminates their well-known toxic actions on the heart. The current study adds a novel mechanism that CTS could exert its biased signaling properties through the NKA signal transducer.

Keywords

cardiotonic steroids; Na/K-ATPase; bias; Digitoxigenin; somalin; ouabain

Significance Statement

Although it is now well accepted that NKA has an ion-pumping independent signaling function, it is still debated whether direct and conformation-dependent NKA/effector interaction is a key to this function. Therefore, this investigation is significant in advancing our understanding of the basic biology of NKA-mediated signal transduction and gaining molecular insight into the structural elements that are important for CTS's biased action.

1. Introduction

Cardiotonic Steroids (CTS), including digitalis drugs such as digoxin and ouabain, are specific inhibitors of Na/K-ATPase (NKA) (Lingrel and Kuntzweiler, 1994; Schatzmann, 1953; Skou, 1988). The positive inotropic effect of CTS on myocardial contractility is due to the inhibition of NKA and subsequent increases in intracellular Ca^{2+} (Eisner and Smith, 1992). Like other inotropes, the FDA approved digoxin produces impressive

hemodynamic effects in chronic heart failure (CHF) patients. However, it distinguishes itself from other pure inotrope drugs because it does not increase CHF patients' mortality, most likely due in part to the recently appreciated signaling through cardiac NKA (Li and Xie, 2009; Pierre et al., 2007).

Over the last decade, studies have revealed that NKA has an ion-pumping independent signaling function (Xie Jeffrey et al., 2013). Precisely, binding of CTS to NKA activates protein and lipid kinases and induces the endocytosis of NKA by changing the function of NKA-interacting proteins (Haas et al., 2000; Kometiani et al., 1998; Liu et al., 2000; Xie, 2001). For example, we have demonstrated that the binding of ouabain to the NKA/Src receptor complex results in the transactivation of the epidermal growth factor receptor (EGFR), resulting in activation of the Ras/Raf/MEK/ERK cascade (Haas et al., 2002). Src activation also leads to stimulation of phospholipase C γ (PLC- γ) and subsequent activation of protein kinase C (PKC) isozymes and Inositol trisphosphate (IP3)- mediated Ca^{2+} signaling (Liu et al., 2007; Mohammadi et al., 2001; Yuan et al., 2005). Moreover, ouabain was found to stimulate PI3K, resulting in the endocytosis of NKA (Liu et al., 2004). CTS activation of NKA-mediated signaling represents a novel mechanism for the modulation of cell growth and apoptosis. It is involved in normal cardiac physiology (Xie et al., 1999) and disease progression such as ventricular hypertrophy and fibrosis (Drummond et al., 2018; Drummond et al., 2016; Elkareh et al., 2007). Importantly, CTS activates these signaling events at concentrations at least an order of magnitude lower than that of inotropy induction.

Several endogenous CTS, including ouabain and marinobufagenin, have been identified, and they play an essential role in the regulation of cardiac structure and function at

concentrations that stimulate NKA signaling without affecting cellular pumping capacity (Hamlyn et al., 1998; Hamlyn et al., 1982; Scheiner-Bobis and Schoner, 2001). At the molecular level, we suggested that receptor NKA regulates protein/protein interaction in a conformation-dependent manner (Ye et al., 2013). Structurally CTS shares a steroid core, a varying number of substituents including a sugar moiety, and a 5- or 6-membered lactone ring, each of which has specific influences on the binding properties to NKA (Cornelius et al., 2013a). While all CTS bind to the E2P state of NKA, they induce different NKA structural changes (Laursen et al., 2015; Yatime et al., 2011). As a consequence, the allosteric control by individual CTS may stabilize NKA in a unique conformation, leading to biased activation of a specific signaling pathway. The growing appreciation that G protein-coupled receptors (GPCRs) can mediate physiologically relevant effects via both G protein and non-G protein effectors has prompted the search for ligands that can "bias" downstream signaling in favor of one pathway over another. Interestingly, animal and clinical studies indicate that the biased β -blocker carvedilol may be a better drug than balanced β -blockers, which also emphasizes the therapeutic value of biased ligands. In support of this CTS bias concept, in this study, we tried to identify diverse structures of CTS that exhibit comparable levels of inhibition of NKA and then test whether these compounds exhibit biased signaling properties *in vitro*, and if so, whether biased signaling capacity alters their *in vivo* pharmacology in mice.

2. Materials and Methods

Materials—the CTS digitoxigenin, somalin, periplogenin and periplogenin 3-O-beta-D-cymaroside were isolated from Chinese herbal medicine *Streptocaulon Juventas* (Lour) Merr. The isolation, verification of compound's structures were reported in another

publication and undergoing publication process, and spectroscopic data were given in supplementary materials in section 1. The CTS digoxin, digoxigenin, ouabain, and ouabagenin were obtained from Millipore Sigma (Burlington, Massachusetts, U.S.A). Structural formulas of compounds were shown in table 1. The pig renal proximal tubule cell line, LLC-PK1, was originally from the American Tissue Type Culture Collection, ATCC (Manassas, VA, USA); Cell culture media, fetal bovine serum, and trypsin were purchased from Invitrogen (Grand Island, NY, USA); the ECL super signal kit was purchased from Pierce; phospho-p44/42 MAPK (ERK1/2, #9101) (Thy 202/204), p44/42 MAPK (ERK1/2, #4695)(137F5) rabbit antibody were from cell signaling. Anti-PKC ϵ rabbit polyclonal (c-15, # sc-214-G), goat anti-rabbit (lot # TK274616), and goat anti-mouse secondary antibodies (lot # TI272496) were from Invitrogen (Thermo Fisher Scientific).

Na/K-ATPase Purification and Na/K-ATPase activity assay—the purified Na/K-ATPase was isolated from the outer medulla of pig kidney as described previously (Xie et al., 1996). The concentration of purified pig kidney enzyme (PKE) was determined by the method of Lowry (27) using bovine serum albumin as the protein standard. The ATPase activity of the PKE was determined according to the assay described below. The specific activities of the Na/K-ATPase of various kidney preparations were in the range of 900 to 1100 $\mu\text{mol Pi/mg/h}$, which was more than 95% of the total ATPase activity. The purity of the PKE was verified by SDS-PAGE, which showed only two bands, one at 110 kDa and the other at 55 kDa, indicative of the α - and β -subunits, respectively, after staining with Coomassie blue.

ATPase inhibitory assay was conducted in class tubes with the final reaction volume of 1900 μ L containing the following components: 100 mM NaCl, 20 mM KCl, 1 mM MgCl₂, 1 mM EGTA, 20 mM Tris-HCl, pH 7.4, and 0.5 μ g of purified Na/K-ATPase. After the compounds were added, the mixtures were incubated at 37°C for 15 min, and reaction was initiated by adding 2 mM ATP-Mg²⁺ mixture. Reactions were carried out for 15 min and then stopped by the addition of 900 μ L of ice-cold 8% trichloroacetic acid (TCA). Reaction mixtures were cleared by centrifugation and assayed for released phosphate by using the BIOMOL Green reagent according to the manufacturer's instructions. In addition, control Na/K-ATPase activity was measured in the presence and absence of 1 mM ouabain and taken as 0 and 100%, respectively. Furthermore, 1 mM ouabain and 0.1% DMSO were included in each plate as a positive and vehicle control, respectively. Control experiments showed that ATP hydrolysis catalyzed by the Na/K-ATPase was in linear range within 30 min of incubation under the above experimental conditions.

Cell Culture and ERK Activation Assay—LLCPK1 cells were cultured in Dulbecco's Modified Eagle's Medium (DMEM), supplemented with 10% fetal bovine serum, 100 units/ml penicillin, 100 μ g/ml streptomycin and 1 μ g/ml puromycin. The cultured cells were routinely tested for possible mycoplasma contamination. When cells reached 90-95% confluence, they were serum-starved overnight and used for experiments. In studies using ouabain and other CTS compounds, duration of exposure for concentration dependent curve was 10 min for ERK activation in all experiments.

PKC ϵ Translocation Assay—LLCPK1 cells were cultured in complete DMEM (containing 10% fetal bovine serum, 100 units/ml penicillin, 100 μ g/ml streptomycin and 1 μ g/ml puromycin). When cells reached 90-95% confluence, they were serum-starved

overnight. Cell lysates were homogenized in buffer A containing (in mmol/L) EGTA(10), EDTA(1), DTT(0.5), PMSF(1), proteinase inhibitor cocktail, Tris-HCl(20), pH 7.5. Homogenates were centrifuged at 100,000 \times g for 1 h at 4 °C. The supernatant designated as the cytosolic fraction was removed and saved. The pellet was sonicated and centrifuged at 25,000 \times g in buffer A containing 1% Triton. The supernatant was collected as the particulate fraction, as we previously described(Duan et al., 2018). Cytosolic and particulate fractions were used in SDS-PAGE and immunoblotting using the anti-PKC ϵ antibody C-15 (Santa Cruz, CA, USA).

Western Blot Analysis—The cells were washed with phosphate-buffered saline and solubilized in ice-cold radioimmune precipitation assay buffer containing 1% Nonidet P-40, 1% sodium deoxycholate, 150 mM NaCl, 1 mM EDTA, 1 mM phenylmethylsulfonyl fluoride, 1 mM sodium orthovanadate, 1 mM NaF, 10 μ g/ml aprotinin, 10 μ g/ml leupeptin, and 50 mM Tris-HCl, pH 7.4, as previously described (15). The cell lysates were then centrifuged at 14,000 rpm, and the supernatants were used for protein assay and subjected to Western blot analysis. The samples were separated on SDS-PAGE (30 μ g/lane) and transferred to a cellulose membrane. The membranes were blocked with a mixture of 1% nonfat dried milk and 1% BSA in TBS-T (10 mM Tris-HCl, 150 Mm NaCl, 0.05% Tween 20, pH 8.0) for one h at room temperature and probed with specific antibodies. The protein signals were detected using an ECL kit, and relative quantitation of protein levels were normalized to total protein loading of signal intensities using ImageJ (NIH).

Cell Viability Assay—To assess the response of LLC-PK1 cells to CTS treatment and stimuli, we utilized cell viability assay to measure the number of live cells in a population.

We used trypan blue exclusion method to distinguish viable cells from nonviable cells. The cells were sub-cultured in 12-well plates at 50,000 cells/well, serum-starved for overnight, and then treated with different concentrations of compounds for indicated time periods. Three wells of control or treated cells were trypsinized and counted manually using a hemocytometer. For distinguishing between dead and viable cells, the samples were diluted with Trypan blue. So that while viewing under a microscope dead cells would appear dark blue.

The quantitation of endocytosed NKA $\alpha 1$ —To analyze NKA endocytosis, we used a YFP- $\alpha 1$ -TCN cell line generated from TCN23-19 cells by first knocking down endogenous NKA, then rescuing NKA expression with YFP-tagged rat $\alpha 1$, which was developed in our lab and reported previously (Cai et al., 2008; Chen et al., 2011; Xu et al., 2021). To test the effect of different structural CTS on the endocytosis of NKA $\alpha 1$, YFP- $\alpha 1$ -TCN cells on glass coverslips were treated with indicated concentrations of CTS for 6 hours and were fixed with 100% prechilled methanol for 30 min at room temperature and washed twice with PBS solution. The coverslips were then mounted. The number of endocytosed NKA $\alpha 1$ was determined in random sections from images captured by fluorescent microscopy acquired at the same magnification. More than 40 cells were examined for each section. For semi-quantitative analysis of the accumulation of NKA in the intracellular compartment, fluorescence intensity of total cell area (Ft) and the intracellular region (Fi) were measured by ImageJ, the percentage of endocytosed NKA $\alpha 1$ was calculated by (Ft-Fi)/Ft. All the data points in the control groups were rescaled to 100%. Values are mean \pm SD from 40 cells in at least three independent experiments.

Labeling of cell surface Na/K-ATPase by biotinylation—Cell surface protein biotinylation was performed as described before (Liu et al., 2005). Briefly, control and treated LLC-PK1 cells were washed twice with ice-cold phosphate-buffered saline (PBS)-Ca-Mg (PBS containing 100 μ mol/L CaCl₂ and 1 mmol/L MgCl₂). Cells were then biotinylated with NHS-ss-Biotin at working concentration of 1.5 mg/ml. Cells were then washed with PBS-Ca-Mg-100 mM glycine for 25 min at 4 degree. Collected cell lysate with lysis buffer and add streptavidin beads to each supernatant. Incubate overnight (12-16 hrs) at 4 degree with end-over-end rotation. Proteins bound to the ImmunoPure™ immobilized streptavidin-agarose beads were eluted and then resolved on sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) followed by immunoblotting.

The Quantification of CTS Bias in NKA α 1 signaling—Base on the given assay in NKA signaling (ERK activation vs NKA endocytosis), using two general approaches to identify CTS-induced signaling bias, which are being validated in GPCRs (Rajagopal et al., 2011). For using the originally proposed qualitative approach refer to as "equimolar comparison" (Gregory et al., 2010a; Gregory et al., 2010b; Rajagopal et al., 2011), we first collected concentration-curve from ERK activation pathway and NKA α 1 endocytosis pathway (figure 2D and figure 3). The responses of these two different assays at the same concentration of ouabain, somalin, or digitoxigenin are then plotted against each other. This curve's shape is a direct comparison of the signaling through the ERK activation and NKA endocytosis. To transform the cell surface NKA α 1 to endocytosed NKA α 1, three steps were taken: 1) we first performed and quantitated the ratio of endocytosed NKA over surface NKA for non-treated LLC-PK1 cells by running

biotinylation assay, ouabain as a positive control. The ratio of endocytosed NKA α 1 (endoNKA) over cell surface NKA α 1 (SurfaceNKA) was obtained as 1.306129 in non-treated (con) cells (EndoNKAcon/SurfaceNKAcon=1.306129, see supplemental section 2), which is also referenced with our previously reported work(Liu et al., 2004; Liu et al., 2005); 2) we then quantitated the ratio of cell surface NKA α 1 (SurfaceNKA) in CTS-treated group over non-treated group (SurfaceNKAtreat/SurfaceNKAcon) from Fig 2C and D. As the total NKA is constantly unchanged, the increased endoNKA in CTS-treated group is established as equation 1; 3) The percentage of endoNKA in CTS-treated groups (Fig 5A X-axis) were then calculated as equation2. From step 1, the endoNKAcon=1.306129*SurfaceNKAcon, the percentage of endoNKA in CTS-treated groups (Fig 5A X-axis) were then finally calculated as equation 3:

Equation1:

$$\text{increased endoNKAtreat} = \text{SurfaceNKAcon} - \text{SurfaceNKAtreat}$$

Equation2:

$$\begin{aligned} \% \text{endoNKAtreat} &= \frac{\text{endoNKAtreat}}{\text{endoNKAcon}} = \frac{(\text{endoNKAcon} + \text{increased endoNKAtreat})}{\text{endoNKAcon}} \\ &= (\text{endoNKAcon} + \text{SurfaceNKAcon} - \text{SurfaceNKAtreat})/\text{endoNKAcon} \\ &= 1 + (\text{SurfaceNKAcon} - \text{SurfaceNKAtreat})/\text{endoNKAcon} \end{aligned}$$

Equation3:

$\%endoNKAtreat$

$$= 1 + (SurfaceNKAcn - SurfaceNKAtreat)/1.306129$$

$$\times SurfaceNKAcn$$

$$= 1 + \left(1 - \left(\frac{SurfaceNKAtreat}{SurfaceNKAcn} \right) \right) / 1.306129$$

Therefore, "endocytosed NKA α 1" used in the calculation of "NKA endocytosis" in Figure 5A was calculated from the endocytosed NKA α 1 in Fig 2C and D using equation 3. The second approach is the quantification of bias factors (equation 4) by using equiactive model used by Robert J. Lefkowitz and coworkers (Rajagopal et al., 2011). In this approach, the doses of ligand (lig) required for an equiactive response for pathway 1 (ERK activation, P1) and 2 (NKA α 1 endocytosis, P2) are extrapolated from fits of each concentration-response curve (figure 2D and figure 3). A bias factor, denoted β , is calculated as the logarithm of the ratio of relative intrinsic activities for a ligand at two different assays compared with a reference agonist. In our case, this comparison was performed using formula (1), which can be calculated from the maximal response (E_{max}) obtained from experiment and the concentration of CTS that provokes a half response (EC_{50}) calculated on Prism (GraphPad). The bias factor quantifies the relative stabilization of ERK activation over NKA α 1 endocytosis compared with endogenous ouabain which is reference ligand (ref).

Equation 4:

$$\beta = \log \left(\left(\frac{E_{max, P1}}{EC_{50, P1}} \frac{EC_{50, P2}}{E_{max, P2}} \right)_{lig} \times \left(\frac{E_{max, P2}}{EC_{50, P2}} \frac{EC_{50, P1}}{E_{max, P1}} \right)_{ref} \right) \quad (1)$$

Statistical Analyses—For NKA inhibition, calculation of IC₅₀ was determined by log(concentration) vs. response curves using the nonlinear iterative three parameter concentration-response curve. Curve fitting was performed with GraphPad Prism 7 (GraphPad Software Inc., San Diego, CA). The quantitative data were subjected to statistical analyses also using GraphPad and are presented as mean ± SD. for all studies of at least 3 times repeat. The differences between groups were determined by ANOVA (multiple groups) followed by multiple comparisons test. Student t-tests were performed to compare two groups. The value of $p < 0.05$ was considered to be significant differences between groups. The nature of the experiments is exploratory, and as such, are not testing a prespecified null hypothesis. P values are, therefore, meant to be descriptive in their interpretation. The number of experimental repetitions was either predetermined based on the level of variation observed in previous work or combined with accumulated data that generated as positive control, and each experiment was reproduced for the number of times as indicated.

3. Results

3.1 Cardiotonic steroids with similar inhibitory effects on purified NKA exhibit different patterns of activation of NKA signaling.

To test the possible biased activity of CTS, we first measured ouabain-sensitive ATPase activities of four pairs of CTS aglycones and glycosides, followed by cell viability assays. As shown in Table 1, CTS exhibited a large difference in their ability to inhibit Na/K-ATPase as previously reported (Stefan Paula, 2005). For example, whereas the half-maximal inhibitory concentration (IC₅₀) between somalin and ouabagenin different by about 50 fold, the IC₅₀ for digitoxigenin and somalin are similar: 0.15 ± 0.04 and

0.18±0.04 μ M, respectively, which is about 5-fold more potent than ouabain. Second, we measured the effect of diverse CTS on renal epithelial (LLC-PK1) cell viability at 100 nM. As shown in Table 1, two glycosides periplogenin 3-O-beta-D-cymaroside and somalin, which share the same cymarose sugar moiety, exhibited more cell toxicity. Unlike somalin, which was cytotoxic, digitoxigenin promotes cell growth at 100 nM, where ouabain, an endogenous NKA ligand, did not affect cell activity. These data indicate a disconnection between the IC₅₀ of CTS in NKA inhibition to their biological effects on cells, suggesting the possible biased biological action of CTS.

To gain further evidence of biased action of CTS, we compared the effects of diverse CTS on ERK activation at 10, 100 nM. Surprisingly, unlike the other CTS, which activates ERK in a concentration-response manner, periplogenin 3-O-beta-D-cymaroside and somalin at these concentrations failed to activate ERK. To corroborate, we utilized a YFP- α 1-TCN cell line, which was generated by first knocking down endogenous NKA then rescuing the NKA knock-down TCN cells with YFP-tagged rat α 1 as a semi-quantitative screening assay for identifying CTS that stimulates the endocytosis of NKA in live cells. As shown in Table 1, all glycosides, including somalin, periplogenin 3-O-beta-D-cymaroside, and digoxin, induced more NKA α 1 endocytosis compared with aglycones at 100 μ M.

We found that digitoxigenin and somalin caused comparable levels of NKA inhibition. However, at a concentration of 100 nM, digitoxigenin stimulated cell proliferation, somalin inhibited cell growth by 30%. These data have shown that ouabain stimulates both protein kinases and NKA endocytosis, while digitoxigenin and somalin bias to ERK activation and endocytosis, respectively.

3.2 CTS regulate different patterns of cell viability through Na/K-ATPase signaling

To understand the molecular mechanism of these opposing effects of somalin and digitoxigenin, we first compared the concentration-response on cell growth in LLC-PK1 cells. Cell numbers were counted in the presence or absence of the indicating concentration of somalin and digitoxigenin, with ouabain as a positive control. As depicted in Figure 1, 2-30 nM ouabain showed a stimulatory effect on cell growth, consistent with what we have reported before (Tian et al., 2009). Like ouabain, 25-100 nM digitoxigenin showed a stimulatory effect on cell growth, and 200-3000 nM caused cell proliferation inhibition in a concentration-dependent manner. In contrast, somalin only induced an inhibitory effect in a concentration-dependent way. As a result, digitoxigenin and ouabain have bell-shaped concentration-response curves, with increased cell viability at low concentrations and decreased cell viability at high concentrations. Compared to ouabain, digitoxigenin has a wider stimulation window, although with the same pattern as ouabain. Structurally, somalin is a glycoside and shares the same aglycone moiety with digitoxigenin. Although similar in structure, with comparable NKA inhibition ($IC_{50} = 0.18 \pm 0.04$, 0.15 ± 0.04 , respectively, in Table 1), they regulate different patterns of cell viability.

3.3 CTS activate biased Na/K-ATPase signal transduction

Ouabain stimulated signal transduction through NKA has been characterized in LLC-PK1 cells, a cell model used extensively to characterize NKA-mediated signal transduction (Liu et al., 2004; Liu et al., 2005; Liu et al., 2002; Tian et al., 2009). To determine whether CTS exhibit biased signaling, we compared digitoxigenin and somalin's impacts on these pathways, using endogenous ouabain as a reference in

LLC-PK1 cells. To analyze NKA endocytosis, we used two complementary assays. First, we generated a YFP- α 1-TCN cell line from TCN cells by first knocking down endogenous NKA, then rescuing NKA expression with YFP-tagged rat α 1. Using fluorescent microscopy, we observed that treatment with 100 μ M of ouabain reduced the surface expression of NKA and consequently increased the accumulation of NKA in the intracellular compartments (Figure 2A). Digitoxigenin induced less endocytosis of NKA, while somalin greatly increased endocytosis, as exhibited by a large accumulation of NKA in the intracellular compartments even at 0.1 μ M. A semi-quantitative analysis (Figure 2B) of NKA in the intracellular compartment revealed that treatment with 100 μ M digitoxigenin induced 50% than the same concentration of ouabain. In contrast, treatment with 0.1 μ M of somalin stimulated a similar level of NKA endocytosis as 100 μ M of ouabain in live cells. To complement the approach, we have developed a quantitative surface biotinylation assay, allowing us to conduct concentration-response studies, as described under "Materials and Methods". As shown in figure 2C and 2D, although all CTS induced Na/K-ATPase α 1 endocytosis in a concentration-dependent manner, they exhibited different potencies. Ouabain induced a balanced effect compared with somalin and digitoxigenin. Somalin was far more potent than digitoxigenin in the case of NKA endocytosis. Endocytosed Na/K-ATPase α 1 was calculated quantitatively from three to five independent experiments for each concentration in figure 2D.

In addition to stimulating endocytosis of NKA, ouabain has been shown to activate ERK with Src-dependent transactivation of EGFR and subsequent activation of Ras/Raf/MEK/ERK cascade (Haas et al., 2000). To evaluate the activation of these

pathways by different CTS, we focused on the activation of ERK in LLC-PK1 cells treated with ouabain, somalin, and digitoxigenin. As shown in figure 3A, unlike ouabain and digitoxigenin, somalin did not activate ERK. In contrast, digitoxigenin displayed a more potent effect on ERK activation than ouabain.

To verify the biased effects of digitoxigenin and somalin are specific through NKA signaling, we performed ERK activation assay in NKA α 1 subunit knockdown and knock-in cell lines that have generated in our lab(Liang et al., 2006) including PY-17 cells, a cell line contain only 8% of the NKA α 1 subunit, and AAC-19 cells, a cell line that were rescued by a rat NKA α 1 for the knockdown PY-17 cells. Functional studies have indicated that knock-down of NKA α 1 decreased the interaction between the Na/K-ATPase and Src and resulted in an increase in basal Src activity. Concomitantly, it also abolished CTS ouabain-induced activation of Src and the downstream ERK cascade(Liang et al., 2006). Based on these, we tested the effect of somalin and digitoxigenin on ERK activation in PY-17 cells and AAC-19 cells. As expected, in figure 3B, both somalin and digitoxigenin failed to activate ERK in PY-17 cells, the NKA α 1 knockdown abolished digitoxigenin-induced activation of ERK. When the knockdown cells were rescued by a rat α 1 (figure 3C), digitoxigenin-induced ERK activation was restored. As the same in LLC-PK1 cells , somalin has no effect on ERK in AAC-19 cells. In addition, digitoxigenin was able to stimulate ERK in the rescued cells at a much higher concentration, which is consistent with the established differences in ouabain sensitivity between pig (LLC-PK1 cells) and rat α 1(AAC-19 cells).

Taken together, these data suggest that digitoxigenin favor protein kinase activation while somalin favor NKA endocytosis without impacting protein kinases activation

(Figures 2-3). It is essential to point out that both digitoxigenin and somalin are about 5-fold more potent than ouabain as NKA inhibitors (Table 1). Yet, they showed distinct concentration-responses in stimulating signal transduction. Significantly, they can produce these signaling events at doses ten times lower than their IC₅₀, meaning that they can initiate signaling without affecting the cellular pumping capacity.

We have shown that ouabain not only activates ERK by Src-dependent transactivation of EGFR and subsequent assembly of Ras/Raf/MEK/ERK cascade but also activates PKC ϵ by Src/EGFR-mediated activation of PLC- γ (Mohammadi et al., 2001). To further test the CTS bias, we also tested the effect of CTS on PKC ϵ activation. As shown in figure 4, LLC-PK1 cells were treated with the indicated concentration of CTS for 15 min. Cytosolic (Cy) and particulate (Pa) fractions were prepared from cell lysates. As expected, ouabain increased Pa/Cy ratios of PKC ϵ contents. Digitoxigenin showed a more potent effect than ouabain, whereas Somalin has no effect on PKC ϵ activation.

CTS-induced endocytosis of NKA and activation of ERK represent a good spectrum of signal transduction to identify the biased effect of CTS. We used two general approaches to quantify CTS-induced signaling bias which is being validated in GPCRs (Rajagopal et al., 2011). The first is an originally proposed qualitative approach refer to as "equimolar comparison" (Gregory et al., 2010a; Gregory et al., 2010b; Rajagopal et al., 2011).

Concentration curve for ouabain, somalin, and digitoxigenin are collected in two assays, including ERK activation assay and NKA α 1 endocytosis assay from figure 2D and figure 3. At the same concentration, the responses of these two assays of ouabain, somalin, and digitoxigenin are then plotted against each other. As shown in figure 5A,

compared with reference balanced ouabain, the shape of the equimolar curve for somalin was biased to NKA α 1 endocytosis but was biased to ERK activation for digitoxigenin. The second approach is the quantification of bias factors by using equiactive model. In this approach, a bias factor, denoted β , is calculated as the logarithm of the ratio of relative activities for somalin and digitoxigenin at two assays, including ERK activation assay and NKA α 1 endocytosis assay, compared with reference ouabain. This comparison was performed using formula (1) as described in "Materials and Methods", which was calculated from the maximal response (E_{max}) obtained from the experiment and the concentration of CTS that provokes a half response (EC_{50}) calculated on Prism (GraphPad). As shown in figure 5B, bias factors driven by ERK activation assay and biotinylation assay demonstrated the biased effect of digitoxigenin and somalin. Somalin relatively stabilized NKA α 1 endocytosis over ERK activation, and digitoxigenin stabilized ERK activation over NKA α 1 endocytosis compared with endogenous ouabain.

4. Discussion

A ligand's ability to stabilize a unique subset of receptor conformations that promotes signaling through distinct pathways to the exclusion of others has been termed ligand bias (Kenakin and Christopoulos, 2013). Ligand bias is associated with the structure of the ligand and receptor systems. After proposing the signaling function of NKA, numerous studies indicate that the NKA is a signal transducer that interacts with several signaling partners (Cui and Xie, 2017; Xie, 2003; Xie and Xie, 2005). CTS as NKA ligands have been shown to regulate protein interactions, and consequently to activate cellular signaling events. This work accumulated evidence of CTS bias that differentially

influences NKA receptor behavior and, consequently, transduces signaling through distinct pathways.

The existence of conformational states of NKA and NKA receptor signaling components lay a fundamental basis for signaling bias. According to the concept of ligand bias proposed in the G-protein coupled receptor (GPCR), CTS bias's eligibility could include CTS-triggered conformational changes and the imposition of altered CTS bound-NKA receptor behavior, which are initiated by selectively starting a biochemical reaction over others. Evidence showing the binding of different CTS could lead to different structural changes in NKA that came from the studies by Klimanova in 2015(Klimanova et al., 2015). It showed variations in fluorescence level upon binding of CTS ouabain or marinobufagenin, using different fluorescent labels targeted to specific residues in the nucleotide-binding domain of NKA, giving direct evidence of distinguished NKA structural changes promoted by different CTS(Klimanova et al., 2015). Also, Azalim and coworkers have confirmed that CTS binding to the NKA is affected by different conformational states of the enzyme in LLC-PK1 cells(Azalim et al., 2020).

Moreover, few works have suggested that even though the signaling pathways are similar, different effects may be elicited by specific CTS in the same measurement system. For example, it has been demonstrated that ouabain attenuates the cardiotoxicity induced by other CTS like digoxin and bufalin(Nesher et al., 2010). This effect is not supported by NKA's ion transport function since ouabain has not been demonstrated to antagonize digoxin-induced inhibition of NKA activity(Nesher et al., 2010).

When considering the possible molecular mechanisms, the biased effect of CTS, evidenced by this work, could be responsible for these functional diversities, which lies in: 1) an initial association of CTS with NKA. Note that the NKA pre-binding conformation(E1P/E2P) are ambiguous in a purified enzyme system (Yoda and Yoda, 1982a; Yoda and Yoda, 1982b), which is a possible implication for physiological ambiguous NKA conformations for CTS binding(Stefan Paula, 2005) that regulated by protein compositions; 2) the conformational rearrangement of the NKA allowing for CTS binding in the binding pocket (Weigand et al., 2014); 3) CTS-stabilized different NKA receptor active states that go on to interact with cellular signaling proteins and lipid to form varieties of complexes in varying quantities; CTS-specific conformation affects direct NKA/effector interaction and the level of protein kinase recruitment. The outcomes of these selective interactions including signal transduction, endocytosis of NKA receptor and formation of an intracellular scaffold that can be evidences for CTS bias; and 4) dissociation process is solely dependent on the nature of the CTS structure. For instance, the sugar moiety, and alternation of the steroid moiety of CTS were reported to influence the dissociation process(Cornelius et al., 2013b; Yoda, 1974). Simple switches in whichever steps could trigger allosteric NKA microprocessor.

By viewing NKA receptor systems, including NKA conformation and the current knowledge about the NKA receptor signaling components in which, direct protein interactions is the molecular mechanism of signal transduction. This work provided evidence that different CTS could trigger biased signaling responses without compromising cells' ion-pumping capacity, presenting a unique cellular response. Before our appreciation of NKA as an essential signal transducer, we have only viewed

CTS as pump inhibitors. Therefore, we have not fully appreciated CTS's potential effects on cardiac structure and remodeling by affecting cells. Recent studies with transgenic animals and using tools that either increase or block NKA-mediated signal transduction have generated a large body of evidence for an important role of NKA in the regulation of cardiac remodeling (Elkareh et al., 2007; Kometiani et al., 2000; Vetteth et al., 2012; Wansapura et al., 2010; Xie et al., 1999). However, although it is now well accepted that NKA has an ion-pumping independent signaling function, it is still debated whether direct and conformation-dependent NKA/effector interaction is a key to this function. Therefore, this investigation is significant in advancing our understanding of the basic biology of NKA-mediated signal transduction and gaining molecular insight into the structural elements that are important for CTS's biased action. It also will lay a solid foundation for the eventual identification of a novel subclass of CTS that is as effective as inotrope but less toxicity than digoxin for congested heart failure patients.

Declaration of competing interest

The authors declare no competing interest.

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Authorship Contributions

Dr. Zijian Xie, Feng Feng, Joseph, I. Shapiro, and Liquan Cai participated in the research design or revision of the manuscript. Dr. Yunhui Xu participated in research design, conducted experiments, performed data analysis and contributed to the writing of the manuscript. Dr. Pauline Marck conducted experiments. Minqi Huang, Dr. Pauline Marck, Dr. Tong Wang and Dr. Jeffrey X. Xie performed data analysis.

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Footnotes

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[†] the author is deceased, January 17th, 2020.

Figure Legends

Figure 1, CTS-effects on cell viability. A, LLC-PK1 cells were sub-cultured in 12-well plates (50,000 cells/well) and serum-starved, three wells of control (Con), and CTS-treated cells were trypsinized and counted after CTS treating for 48 hours. The quantification data are from three to four independent experiments. The individual data points are mean \pm SD. **, $p < 0.01$ compared with 100 nM of ouabain, unpaired t-test. B, after digitoxigenin (Digt) treatment, three wells of control (Cont) and Digt-treated cells were trypsinized and counted at indicated time points. The same experiments were repeated three times. The individual data points are mean \pm SD. *, $p < 0.05$ compared with vehicle Cont, unpaired t-test.

Figure 2, CTS-effects on Na/K-ATPase $\alpha 1$ endocytosis. A, TCN-YFP- $\alpha 1$ cells were treated with CTS for 6 hours and imaged under a fluorescence microscope. B, semi-quantitative endocytosed NKA $\alpha 1$ was calculated by counting the accumulation of NKA in the intracellular compartments. Data were quantified from four independent experiments and standardized to vehicle control. Individual values are presented as mean \pm SD. Data was analyzed with one-way ANOVA followed by Sidak's multiple

comparisons test to determine statistical significance. **, $P < 0.01$, ***, $P < 0.001$, ****, $P < 0.0001$. C, cell surface NKA $\alpha 1$ was measured by biotinylation assay described under “Materials and Methods”. Protein was collected after treatment with different concentrations of CTS. Cell lysates were separated by SDS-PAGE and analyzed by Western blot for NKA $\alpha 1$. D, quantitative dose-response endocytosed NKA $\alpha 1$ was calculated from three to five independent experiments for each concentration. The individual data points are mean \pm SD.

Figure 3, CTS-effects on ERK activation in LLC-PK1 cells. Cells were treated with indicated concentration of CTS, probe for active ERK by Weston blot. Upper panel: representative western blot. Lower panel: dose curve for respective CTS treatment presented as mean \pm SD, $n=6$ for ouabain (Oub) group, $n=9$ for somalin (Som) and digitoxigenin (Digt). The EC_{50} for Oub and Digt are 6.52 nM, 95% CI 3.98-13.80 nM, and 3.85 nM, 95% CI 1.96-7.408 nM, respectively.

Figure 4, CTS-effects on ERK activation in PY17 and AAC-19 cells. Cells (A, PY17. B, AAC-19) were treated with indicated concentration of CTS for 10 minutes and probe for phospho-ERK by Weston blot. Upper panel: representative western blot. Lower panel: grouped table format for respective CTS treatment presented as mean \pm SD, $n=3$. C, AAC-19 cells were treated with indicated concentration of CTS for 10 minutes and probe for phospho-ERK by Weston blot. Upper panel: representative western blot. Lower panel: grouped table format for respective CTS treatment presented as mean \pm SD, $n=3$. Using a grouped analyses of multiple t tests-one per row. The statistical significance determined using the Holm-Sidak method, with $\alpha=0.05$. Each row was analyzed individually, without assuming a consistent SD.

Figure 5, CTS-effects on PKC ϵ activation in LLC-PK1 cells. LLC-PK1 cells were treated with indicated CTS for 15 min. Cytosolic (Cy) and particulate (Pa) fractions were prepared from cell lysates and PKC ϵ activation as measured by Pa/Cy ratios of PKC ϵ contents were compared by Weston blot analysis. Left panel: representative blots are shown. Right panel: dose curve for respective CTS treatment. Individual values are presented as mean \pm SD, n=3-6 for ouabain (Oub) group, n=4-10 for somalin (Som), and n=4-12 for digitoxigenin (Digt) group. The n numbers are varies because some of the data are combined from different repeat with different concentration sets by different person.

Figure 6, identification of CTS-induced bias in NKA α 1 signaling. A, an equimolar comparison between the ERK activation and the endocytosis of NKA α 1 demonstrate that Som is dramatically biased to NKA endocytosis, and Digt is slightly biased to ERK activation. B, bias factors were calculated by using the equiactive model as described under "Materials and Methods". Equiactive comparison between ERK activation and NKA α 1 endocytosis demonstrate biased effect of Digt and Som. Data are presented as mean \pm SD. **, p<0.01 by t-test.

Table

Table 1, chemical structures of CTS and effects of CTS on Na/K-ATPase inhibition, cell growth rate, ERK activation, and Na/K-ATPase α 1 endocytosis.

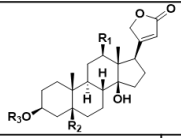
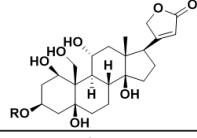
Name	Digitoxigenin-based compounds							
	R1	R2	R3	NKA inhibition	Cell growth rate (100% of non-treated)	ERK activation (100% of non-treated)		NKA α 1 endocytosis (100% of non-treated)
				IC ₅₀ , 95% CI, μ M	100 nM	10 nM	100 nM	100 μ M
Digitoxigenin	H	H	OH	0.15, 0.11-0.19	131.64 \pm 0.07	210.43 \pm 27.75	276.39 \pm 33.55	392.33 \pm 144.64
Somalin	H	H	cymarose	0.18, 0.12-0.24	69.79 \pm 0.56	95.00 \pm 12.72	117.66 \pm 12.35	820.25 \pm 51.98
Periplogenin	H	OH	OH	0.31, 0.24-0.54	116.03 \pm 0.83	141.84 \pm 12.41	156.89 \pm 18.41	141.00 \pm 16.45
periplocymarin	H	OH	cymarose	0.62, 0.38-1.13	31.37 \pm 0.07	92.09 \pm 3.36	99.14 \pm 8.70	920.75 \pm 58.78
Digoxigenin	OH	H	OH	0.49, 0.34-0.69	83.26 \pm 0.13	112.92 \pm 7.10	141.65 \pm 17.01	555.00 \pm 47.26
Digoxin	OH	H	trisdigitoxose	1.78, 1.34-2.43	102.64 \pm 0.26	105.82 \pm 7.10	159.93 \pm 30.52	779.00 \pm 111.38
	Ouabagenin-based compounds							
	R1							
Ouabagenin	H			7.76, 5.73-11.74	115.03 \pm 0.07	103.69 \pm 11.36	129.36 \pm 19.63	510.00 \pm 38.50
Ouabain	rhamnose			1.01, 0.85-1.36	100.00 \pm 0.85	174.22 \pm 17.44	220.43 \pm 19.33	572.5 \pm 36.08

Figure 1

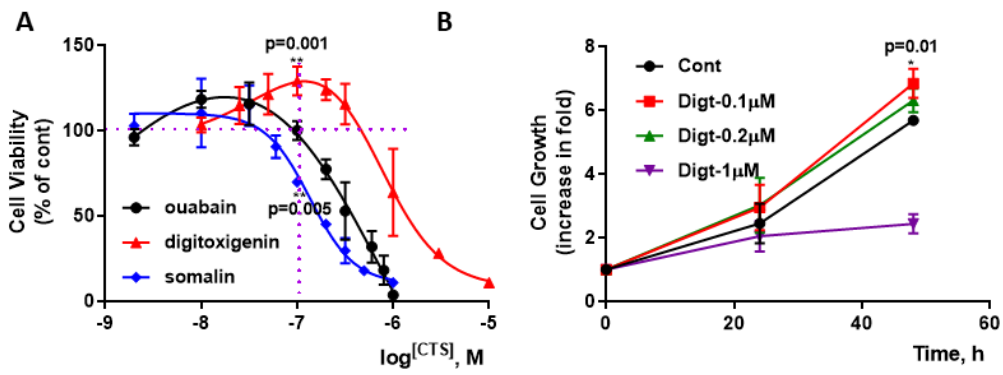


Figure 2

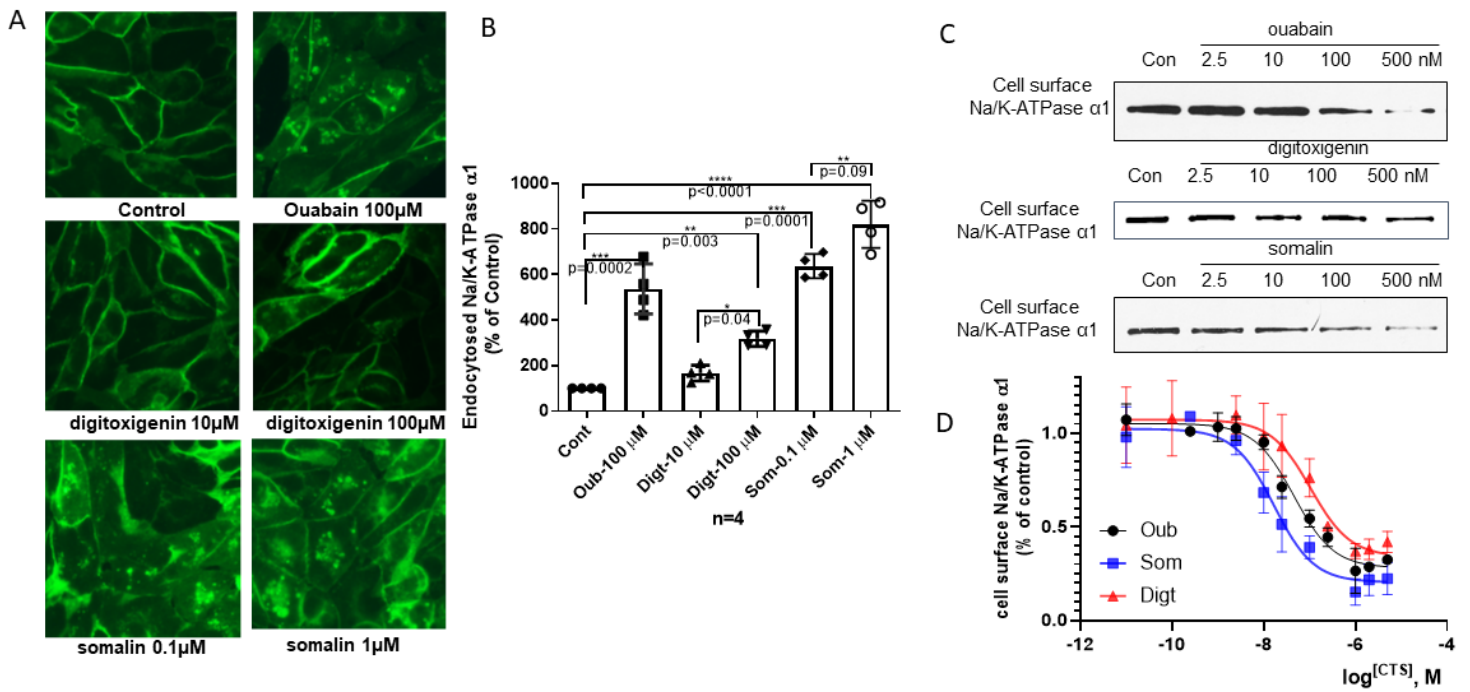


Figure 3

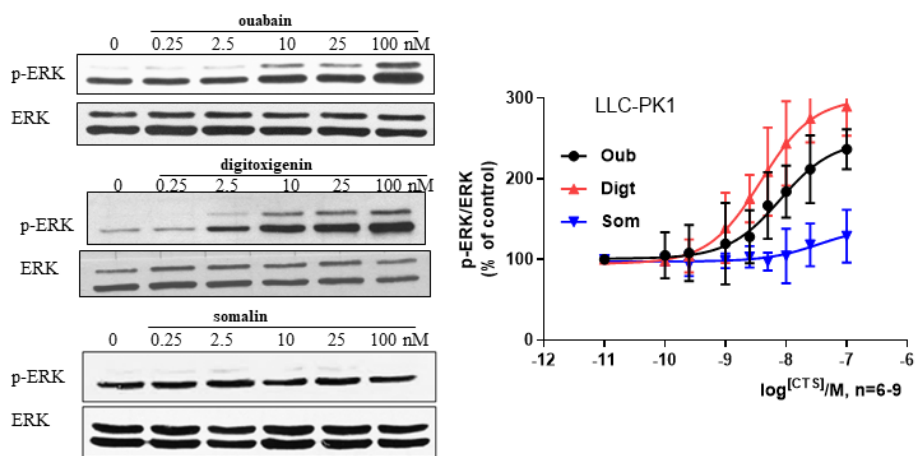


Figure 4

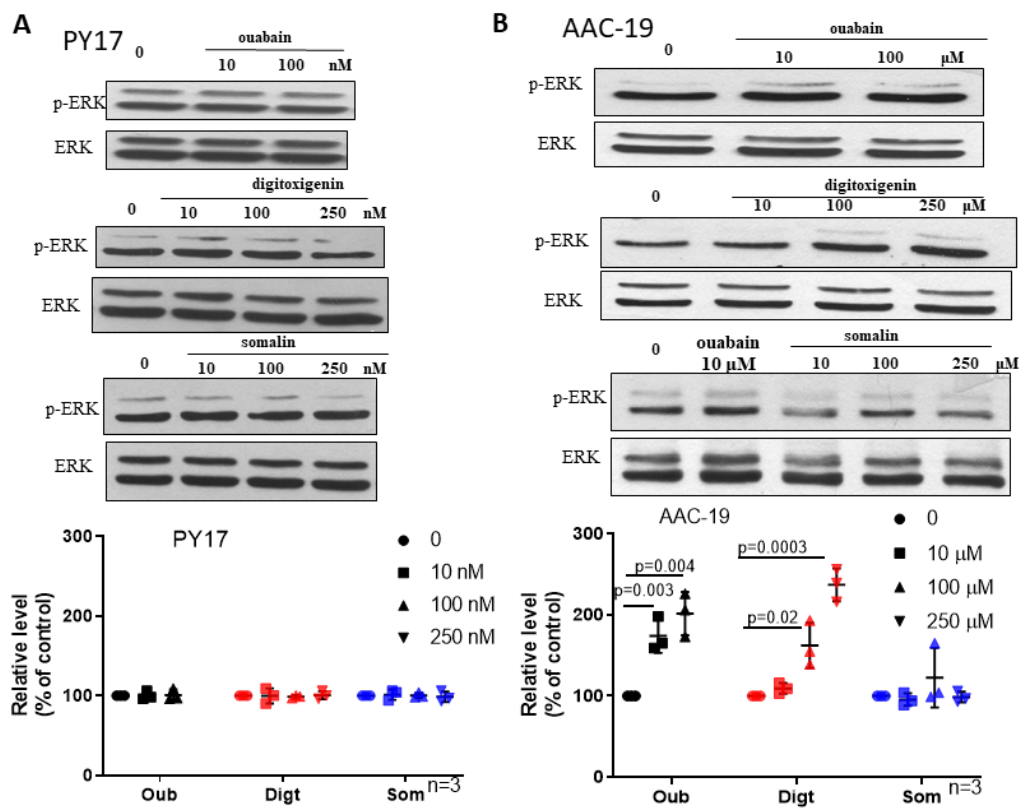


Figure 5

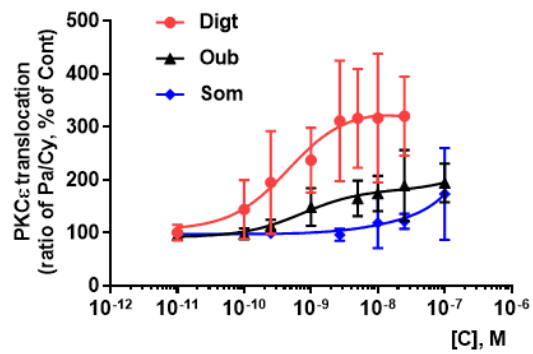
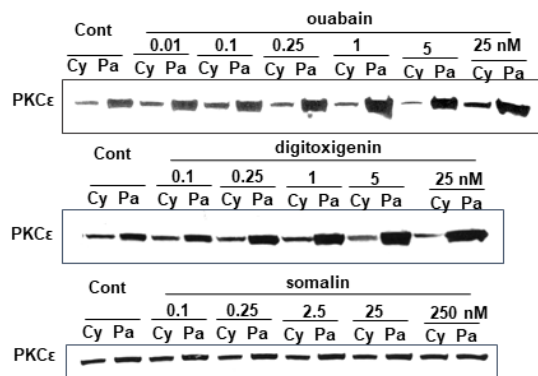


Figure 6

