Cysteine residues 244 and 458-459 within the catalytic subunit of Na,K-ATPase control the enzyme’s hydrolytic and signaling function under hypoxic conditions

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Abstract: Our previous findings suggested that reversible thiol modifications of cysteine residues within the actuator (AD) and nucleotide binding domain (NBD) of the Na,K-ATPase may represent a powerful regulatory mechanism conveying redox- and oxygen-sensitivity of this multifunctional enzyme. S-glutathionylation of Cys244 in the AD and Cys 454-458-459 in the NBD inhibited the enzyme and protected cysteines’ thiol groups from irreversible oxidation under hypoxic conditions. In this study mutagenesis approach was used to assess the role these cysteines play in regulation of the Na,K-ATPase hydrolytic and signaling functions. Several constructs of mouse 1 subunit of the Na,K-ATPase were produced in which Cys244, Cys 454-458-459 or Cys 244-454-458-459 were replaced by alanine. These constructs were expressed in human HEK293 cells. Non-transfected cells and those expressing murine 1 subunit were exposed to hypoxia or treated with oxidized glutathione (GSSG). Both conditions induced inhibition of the wild type Na,K-ATPase. Enzymes containing mutated mouse 1 lacking Cys244 or all four cysteines (Cys 244-454-458-459) were insensitive to hypoxia. Inhibitory effect of GSSG was observed for wild type murine Na,K-ATPase, but was less pronounced in Cys454-458-459Ala mutant and completely absent in the Cys244Ala and Cys 244-454-458-459Ala mutants. In cells, expressing wild type enzyme, ouabain induced activation of Src and Erk kinases under normoxic conditions, whereas under hypoxic conditions this effect was reversed. Cys454-458-459Ala substitution abolished Src kinase activation in response to ouabain treatment, uncoupled Src from Erk signaling, and interfered with O2-sensitivity of Na,K-ATPase signaling function. Moreover, modeling predicted that S-glutathionylation of Cys 458 and 459 should prevent inhibitory binding of Src to NBD. Our data indicate for the first time that cysteine residues within the AD and NBD influence hydrolytic as well as receptor function of the Na,K-ATPase and alter responses of the enzyme to hypoxia or upon treatment with cardiotonic steroids.

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Research paper

Cysteine residues 244 and 458–459 within the catalytic subunit of Na,K-ATPase control the enzyme's hydrolytic and signaling function under hypoxic conditions

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ABSTRACT

Our previous findings suggested that reversible thiol modifications of cysteine residues within the actuator (AD) and nucleotide binding domain (NBD) of the Na,K-ATPase may represent a powerful regulatory mechanism conveying redox- and oxygen-sensitivity of this multifunctional enzyme. S-glutathionylation of Cys244 in the AD and Cys 454-458-459 in the NBD inhibited the enzyme and protected cysteines’ thiol groups from irreversible oxidation under hypoxic conditions. In this study mutagenesis approach was used to assess the role these cysteines play in regulation of the Na,K-ATPase hydrolytic and signaling functions. Several constructs of mouse α1 subunit of the Na,K-ATPase were produced in which Cys244, Cys 454-458-459 or Cys 244-454-458-459 were replaced by alanine. These constructs were expressed in human HEK293 cells. Non-transfected cells and those expressing murine α1 subunit were exposed to hypoxia or treated with oxidized glutathione (GSSG). Both conditions induced inhibition of the wild type Na,K-ATPase. Enzymes containing mutated mouse α1 lacking Cys244 or all four cysteines (Cys 244-454-458-459) were insensitive to hypoxia. Inhibitory effect of GSSG was observed for wild type murine Na,K-ATPase, but was less pronounced in Cys454-458-459Ala mutant and completely absent in the Cys244Ala and Cys 244-454-458-459Ala mutants. In cells, expressing wild type enzyme, ouabain induced activation of Src and Erk kinases under normoxic conditions, whereas under hypoxic conditions this effect was inverted. Cys454-458-459Ala substitution abolished Src kinase activation in response to ouabain treatment, uncoupled Src from Erk signaling, and interfered with O2-sensitivity of Na,K-ATPase signaling function. Moreover, modeling predicted that S-glutathionylation of Cys 458 and 459 should prevent inhibitory binding of Src to NBD. Our data indicate for the first time that cysteine residues within the AD and NBD influence hydrolytic as well as receptor function of the Na,K-ATPase and alter responses of the enzyme to hypoxia or upon treatment with cardiotonic steroids.

1. Introduction

Na,K-ATPase is a ubiquitous enzyme present in plasma membranes of almost all cells. It is formed by a catalytic α and a regulatory β subunits of which several isoforms are known (α1-α4 and β1-β3). In some tissues one more regulatory subunit belonging to the family of FXYD proteins joins the αβ dimers [7,8]. Na,K-ATPase is a major gradient-forming enzyme moving Na+ and K+ against their electrochemical gradients. In addition Na,K-ATPase is a receptor for cardiotonic steroids and a mediator of complex signaling cascades involving kinases, Ca2+ and mitochondrial free radical burst [1,15]. Whereas lower doses of cardiotonic steroids evoke signaling response of the Na,K-ATPase, at higher concentrations these compounds inhibit transport function of the enzyme [56].

Consuming 20–85% of ATP in cells and tissues, Na,K-ATPase is one of the major ATP sinks in all cells. It converts energy of ATP to that of transmembrane Na+ and K+ gradients. The latter is used to sustain brain function and the muscle contractility, nutrients uptake and multiple other processes. Na,K-ATPase is also known to be a sensor of redox state and O2 availability [10,11,55,56]. Decrease in O2 levels...
comprises survival of the organism as the ion gradient maintenance may no longer be effectively preserved due to the quick (and reversible) inactivation of Na,K-ATPase [12,14,16–19]. Furthermore, systemic hypoxia triggers release of low doses of cardio
tonic steroids from adrenal cortex into the circulation. Binding of endo
genous cardioactive steroids to the Na,K-ATPase and activation of the receptor function of the enzyme is associated with initiation of signal
ging cascades involving Src-Erk kinases [18]. Tuning of the Na,K-ATPase role in O2 availability translates into a substantial reduction in ATP consumption under hypoxic conditions, a powerful adaptive response allowing to avoid acute terminal ATP depletion. Complex mechanisms of adjustment of the Na,K-ATPase activity to the changes in environmental oxygen levels are largely coupled to the alterations in redox environment [11]. Whereas suppression of transport and hydro
dlytic function of the Na,K-ATPase under hypoxic conditions is well
described, little is known about the impact of hypoxia on the receptor function of the enzyme.

Oxidative thiol modifications were recently shown to play a decisive role in control of ion transport and ATP hydrolysis by the Na,K-ATPase in hypoxic cells [23,31,54,66]. Onset of hypoxia is associated with a transient free radical burst from the mitochondria along with gradual ATP depletion [26]. Accumulation of oxidized glutathione (GSSG) that follows the oxidative burst joins the reaction of dithiol exchange with regulatory thiols of multiple proteins including Na,K-ATPase. In our previous study we have explored the localization of these regulatory thiols within the catalytic α1 subunit of the Na,K-ATPase [54]. Functional tests as well as mass spectrometry and in silico modeling indicated that regulatory thiols are most likely located within the ATP binding pocket of the catalytic α subunit of the Na,K-ATPase interfering thereby with ATP binding to the nucleotide binding domain (NBD) [54]. Exposure of purified Na,K-ATPase protein to GSSG in concentra
tions comparable with those found in hypoxic tissues resulted in complete suppression of the enzyme’s hydrolytic activity [54]. In the ab
cence of ATP inhibition of the Na,K-ATPase occurred within 5–10 min of treatment, and the activity could be restored by DTT as well as by exposure to glutaredoxin 1/NADPH mixture. Binding of ATP to the Na,K-ATPase made it insensitive to the inhibitory action of GSSG [54]. Induction of S-glutathionylation of the α-subunit and the concomitant decrease in hydrolytic activity of the Na,K-ATPase was confirmed in hypoxic myocardial tissue and several cell lines [43,51,54,66].

Four cysteine residues within the actuator domain (AD) (Cys244) and NBD (Cys 454, 458 and 459) of the α1 subunit were suggested as targets of regulatory S-glutathionylation based on the data of mass
spectrometry and in silico modeling [54]. One of them, Cys454, was later identified as inaccessible to S-glutathionylation after the protein folding and could thus only become S-glutathionylated in the ER [43]. In this study point-mutations approach was applied to evaluate the role of regulatory cysteines in O2-sensitivity of hydrolytic and receptor function of the Na,K-ATPase. Wild type murine catalytic α1 subunit as well as the protein in which the candidate regulatory cysteines in the AD or/and NBD were substituted by alanine were expressed in a human embryonic kidney derived cells (HEK-293). As murine α1β isozyme requires 1000-fold higher concentrations of cardioactive steroid, oua
bain, for inhibition than the human α1β isozyme, the activity of transfected mouse enzyme could be easily distinguished from the human endogenous Na,K-ATPase. Hydrolytic activity of the Na,K-ATPase as well as the signaling cascade triggered by ouabain binding to the enzyme were assessed in normoxic and hypoxic transfected HEK 293 cells. We further assessed sensitivity of wild type and mutated enzyme to GSSG.

The obtained results indicate that Cys244 is of key importance in redox-sensitivity of the Na,K-ATPase activity. Substitution of Cys458 and 459 to alanines compromises hypoxia-sensitivity and receptor function of the enzyme.

2. Materials and methods

2.1. Plasmid preparation

Wild type, murine Na,K-ATPase α1 subunit was subcloned into pcDNA3.1/myc-His (Invitrogen) from IMAGE clone 4951288, posi
tioning the myc epitope and 6×His tag at the C terminus of Na,K-
ATPase α1 subunit. The following site-directed mutagenesis strategy was chosen. Cysteines 244, 454-458,459 and a combination of all four cysteine residues were replaced by alanine and, as pilot experiments showed, resulted in production of a functional protein. Mutagenesis reactions were performed with the QuikChange II site-directed muta
genesis kits (Agilent). Mutagenesis primers were designed using the Agilent online primer design tool.

C244A (forward): GCCTTCTTCTCAACAAAGCTGTGGAAGGAACCCGGC
CCC454/458/459AAA (forward): CAGAGTGCAGGCTTCTAAGAGGC
CATTAAGATGCAGGCGCAGGC

All mutations were confirmed by sequencing the entire ATP1a1 cDNA. Plasmids were multiplied in E. coli and purified for transfection into HEK293 cells.

2.2. Cell culture and transfection

Human embryonic kidney-derived cell line HEK293 was used for transfection with mouse α1 or mock plasmids. Cells were plated in 25 cm2 flasks 24 h prior to the transfection at the confluence of 25–30% and maintained in DMEM supplemented with 20% fetal calf serum, 100 U/ml penicillin, 100 μg/ml streptomycin and 1 mM Na-pyruvate at 37 °C and 5% CO2 in gas phase. Lipofectamine 2000 Transfection Reagent (Thermo Fisher Scientific) was used to transfect HEK293 cells with the plasmids (8 μg per transfection) containing the wild type and mutated mouse Na,K-ATPase α1 subunit. Standard transfection protocol of the producer was used (https://tools.thermofisher.com/content/sfs/manuals/Lipofectamine_2000_Reagent_protocol.pdf). Neomycin (418) was added to the cells 24–48 h after the transfection at concentration of 1000 μg/ml and selection for the transfected cells performed over 72 h. Transfected cells were then directly used for experiments.

2.3. Immunoblotting for the his-tag

Cells were lysed in buffer containing 500 mM NaCl, 25 mM Tris-
HCl, 10 mM imidazol (pH 7.5), 1% Nonidet-P40, 0.1% SDS, 1% sodium deoxycholate and 1 μM PMSF and incubated at 4 °C for 1 h under dis
tinuous stirring. The samples were then centrifuged at 13,000 ×g for 10 min and the supernatant was collected. Separation of His-tagged murine Na,K-ATPase was performed using high performance Ni Sepharose (GE Healthcare). Ni Sepharose was washed in buffer contain
ing 25 mM Tris-HCl, 500 mM NaCl, 10 mM imidazole (pH 7.5). Thereafter 25 μl of Ni Sepharose was mixed with 250 μl of cell lysates and incubated at 4 °C for 1 h under constant stirring. Ni Sepharose with Na,K-ATPase bound to it was sedimented by centrifugation at 16

000×g for 1 min, and supernatant containing released Na,K-ATPase was collected, separated on SDS-PAGE and transferred to a PVDF membrane. After membrane blocking in 5% non-fat milk in PBS the detec
tion of His-tagged Na,K-ATPase was performed using anti-his-tag antibody (dilution 1:1000) and mouse monoclonal anti-Na,K-ATPase α1 antibody clone C464-6 (Upstate, Millipore, dilution 1:10000) and secondary horseradish peroxidase-conjugated antibodies along with enhance chemiluminiscence SuperSignalTM West Femto Maximum Sensitivity Substrate kit (ThermoFisher Scientific). Chemiluminiscence was detected using Bio-Rad ChemiDoc MP imaging system.
2.4. Immunoblotting for S-glutathionylated α1 subunit

S-glutathionylation of Na,K-ATPase α1 subunit was estimated using immunoblotting. Proteins from cell lysates were separated on the SDS-PAGE (without β-mercaptoethanol) and transferred to PVDF membranes. After blocking, mouse monoclonal anti-glutathione antibodies (Chemicon Millipore, MABS310) were applied. Mouse monoclonal anti-α1 antibody clone C464-6 (Upstate, Millipore) was used to detect total amount of the α1 subunit. Secondary horseradish anti-mouse antibodies were used with signal enhancement by chemiluminescence SuperSignal™ West Femto Maximum Sensitivity Substrate kit (ThermoFisher Scientific). Chemiluminescence was detected using Bio-Rad ChemiDoc MP imaging system.

Densitometric analysis was performed using Image Lab (Bio-Rad) program and the results were presented as a ratio of S-glutathionylated fraction of α1 subunit to total amount of α1 subunit in the corresponding lane. This ratio in control non-treated sample was taken as 1.

2.5. Hypoxic exposure and detection of cell viability

Short-term hypoxic exposure was performed using pre-equilibrated hypoxic media that were applied to the cells within hypoxic bench (InVivo400 Ruskind). Longer incubations were performed in hypoxic incubators (Hercoses). Hypoxic conditions were defined as 1% O₂ up to 24 h (long-term viability experiments, or 0.2% O₂ for 30–60 min (acute signaling responses, for details see text and figures legends). The cells were then washed free from the culture medium with phosphate buffer or Tyrode buffer containing (in mM): 145 NaCl, 5 KCl, 2 CaCl₂, 1 MgCl₂, 5 glucose, and 10 HEPES/NaOH, pH 7.4, and de-attached using scrapers or by trypsinization. So obtained, cells and lysates were stored in liquid nitrogen or used immediately for cell viability assessment. ADAM-17 Automatic Cell Counter was used to test for the cell viability and cell number after the exposure of cells to hypoxia 1% O₂. Decrease of viability was assessed by registration of the percent of propidium iodide-positive cells (dead cells) related to the total number of cells. Toxicity of the membrane-permeable analogue of GSSG, tetraethyl-GSSG (ttGSSG), in HEK 293 cells was assessed using MTT assay as described earlier [51]. No changes in cell viability was detected in cells treated with ttGSSG in concentrations up to 2 mM over 24 h (Suppl. Fig. 1).

2.6. Treatment of cells with oxidized glutathione

GSSG and ttGSSG were dissolved in 300 mM HEPES/NaOH buffer (pH 7.4 at room temperature) to obtain stock solution of 10 mM. Cell culture medium in 25 cm² flasks in which HEK293 cells were growing was replaced by 1 ml Tyrode buffer. Oxidized glutathione derivatives were then added to the buffer to reach final concentration of 1 or 0.5 mM for GSSG or ttGSSG respectively. pH of Tyrode did not change after addition of GSSG or ttGSSG. Cells were incubated with ttGSSG for 1 h in the incubator in the atmosphere of air with 5% CO₂ at 37 °C. Treatment with GSSG was performed immediately before the cells were de-attached using scrapers and frozen in liquid N₂. Thus, the membranes of cells were in contact with GSSG upon thawing.

2.7. Detection of hydrolytic activity of Na,K-ATPase

About 1000-fold difference in affinity to ouabain is known to exist between the human and murine Na,K-ATPase [15,48] making it possible to distinguish between the activities of human and murine isoforms. Hydrolytic activity of Na,K-ATPase in cell lysates was measured as ouabain-sensitive phosphate (Pi) production in the medium containing the following saturating levels of ions and substrates (mM): 130 NaCl, 20 KCl, 3 MgCl₂, 3 ATP, 30 HEPES/NaOH (pH 7.4, 37 °C). 1 mM GSSG was additionally present in the medium for cells pre-treated with GSSG and ttGSSG during the activity measurements. Details of the protocol were described elsewhere ([54,57]). 10 μM ouabain was used to selectively block human isoyme whereas 2 mM ouabain was effectivly blocking both human and mouse Na,K-ATPase activity. Activity of murine ATPase could then be calculated by subtracting the activity of human ouabain-sensitive isoymes from the total Na,K-ATPase activity. Enzyme activity was expressed per mg protein per hour and protein content was detected using Bradford method.

2.8. Assessment of Src and Erk1/2 phosphorylation state by immunoblotting

Phosphorylation of Src and Erk1/2 kinases was evaluated in cell lysates of non-transfected and transfected HEK293 cells that were incubated with or without 250 μM ouabain under normoxic (20% O₂) or hypoxic (0.2% O₂) conditions. Incubation at normal and low oxygen levels was performed in Tyrode buffer for 15 min prior to addition of 250 μM ouabain and for further 30 min in the presence of cardiotonic steroid. The cells were then de-attached by scrapers and sedimented by centrifugation for 10 min at 100 × g. Cells were lysed in the RIPA buffer (25 mM tris-HCl, pH 7.6, 150 mM NaCl, 1% Nonidet-P40, 0.1% SDS, 1% sodium deoxycholate) containing 1 μM of PMSF with stirring at 4 °C for 1 h. The probes were then centrifuged at 13,000 × g for 10 min and the supernatant was collected. Proteins of cell lysates were separated on SDS-PAGE and transferred to a nitrocellulose membrane. After membrane blocking in 5% BSA (bovine serum albumin) in PBST, the detection of phospho (Tyr 416) Src or Erk1/2 and total Src or Erk1/2 was carried out by incubating the membrane in the solution of appropriate rabbit polyclonal antibodies (both from Cell Signaling Technology) in PBST. Visualization of the proteins was performed by the appropriate horseradish peroxidase-conjugated secondary antibodies followed by the enhanced luminiscent SuperSignal ™ West Femto Maximum Sensitivity Substrate kit (Thermo Scientific). Chemiluminescence was detected using Bio-Rad ChemiDoc MP instrument. Densitometric analysis was performed with Image Lab program (Bio-Rad) and the results were expressed as ratio of phospho-Src or phospho Erk1/2 to total Src or Erk1/2 band intensity (phospho-Src/Src; phospho-Erk1/2/Erk1/2).

2.9. Modeling of binding of Src kinase to the NBD of Na,K-ATPase

Modeling of the complex was performed using the structure of porcine Na,K-ATPase (PDB id 3wgu) solved at 2.8 Å resolution, and the structure of human Src kinase (PDB id 2src). Docking was performed using Na,K-ATPase NBD (377–588 residues) and kinase domain of Src kinase (residues 267–520). Global docking of these domains has been carried out with SwamDock, GRAMM_X, ZDock and ClusPro servers. Thereafter the full-size protein structures were reconstructed. Qasdom server (http://qasdom.eimb.ru/qasdom.html) was used to identify interactions between the NBD and kinase domain of Src kinase in obtained models.

2.10. Statistical analysis

Statistical module of SigmaPlot v.13 (SigmaPlot) was used for statistical analysis. Rank sum test or signed rank test were used for analysis whenever appropriate. p < 0.05 was considered as significant difference.

3. Results

3.1. HEK293 express wild type and mutated murine α1 subunit forming functional Na,K-ATPase

Several cell lines were generated based on the HEK293 cells that expressed wild type or mutated forms of mouse α1 subunit of Na,K-ATPase along with homologous α1 subunit (α1h), the only isoform expressed in this cell line [13]. Along with endogenous α1h α1
form of Na,K-ATPase transfected cells expressed either mouse wild type enzyme (α1wtβ1) or one of the following three mutated mouse enzymes: Cys244Ala mutant (α1m-1CAβ1), Cys454-458-459Ala mutant (α1m-3CAβ1), or Cys244-454-458-459Ala mutant (α1m-4CAβ1). Expression of the transfected mouse α1 subunits was confirmed by means of immunoblotting against the histidine tag (Suppl. Fig. 2A). Cells expressing any of mouse α1 subunits showed increased levels of the α1 protein compared to the non-transfected HEK293 cells (Suppl. Fig. 2B). Functionality of the compound Na,K-ATPase molecule formed by the transfected mouse α and human β subunits was tested. Ouabain at concentration of 10 μM was used to block human α1hβ1 Na,K-ATPase. Mouse Na,K-ATPase activity was insensitive to 10 μM, but completely blocked by 2 mM ouabain. The enzymes containing mouse α1 subunit were active and contributed by 40–50% into the total hydrolytic activity of the Na,K-ATPase (Suppl. Fig. 2C).

3.2. Replacement of Cys244 by Ala makes the enzyme insensitive to GSSG

Two sets of experiments were performed to address the role of the Cys 244 in the AD and Cys 454, 458 and 459 in the NBD in sensitivity of Na,K-ATPase to GSSG. In the first set of experiments non-transfected HEK293 cells as well as the cells expressing α1wtβ1, α1m-1CAβ1, α1m-3CAβ1 and α1m-4CAβ1 Na,K-ATPase were lysed by freezing-thawing cycle in the presence of 1 mM GSSG. Activity of endogenous human Na,K-ATPase in cell lysates was suppressed after GSSG treatment by 40% (Fig. 1). For the enzyme formed by the wild type mouse α1 and human β1 subunits suppression of activity after GSSG treatment reached 60% (Fig. 2A). Mutants α1m-1CAβ1 and α1m-4CAβ1 were insensitive to the inhibitory action of GSSG, while α1m-3CAβ1 mutant inhibitory effect of GSSG was abolished only partially. These findings were verified in the next set of experiments in which sensitivity of the Na,K-ATPase to GSSG was tested in intact cells. Plasma membrane is known to be impermeable to GSSG and loading of cells with it was performed using its membrane-permeable esterified homologue, tGSSG. Incubation of non-transfected HEK293 cells with tGSSG induced S-glutathionylation of the Na,K-ATPase α1 subunit (Suppl. Fig. 3A). Exposure of transfected cells to tGSSG resulted in a decrease in α1wtβ1 Na,K-ATPase activity by more than 40% (Fig. 2B). Activity of the α1m-1CAβ1 and α1m-4CAβ1 mutants was unaffected by tGSSG (Fig. 2B). S-glutathionylation levels of the untreated mutant-α1 subunits were similar to those in wild type murine α1 subunit (Suppl. Fig. 3B).

Fig. 1. Sensitivity of endogenous human Na,K-ATPase to GSSG and hypoxia. Non-transfected HEK 293 cells were exposed to 1% O2 (hypoxia) for 24 h or to 1 mM GSSG during lysis and Na,K-ATPase activity measurements. Data are means ± SD for 3 experiments. * stands for the p < 0.05 compared to control.

Endogenous human Na,K-ATPase in non-transfected HEK293 cells responded to hypoxia (1% O2, 24 h) with a loss of ~40% activity (Fig. 1). Similar response was observed for the α1mwt-β1 enzyme (Fig. 3A). However, α1m-3CAβ1 mutant was not inactivated by hypoxia. The α1m-1CAβ1 and α1m-4CAβ1 enzymes in which Cys244 was replaced by Ala responded to deoxygenation with an increase in their hydrolytic activity (Fig. 3A). Incubation of non-transfected HEK293 cells and cells with α1mwt-β1 Na,K-ATPase at 1% O2 for 24 h resulted in decrease of cell viability by 20% (Fig. 3B). The viability of cells expressing the α1m-4CAβ1 mutant was lower than that of cells expressing α1mwt-β1 enzyme. Thus, the cells expressing the GSSG-insensitive α1m-4CAβ1 mutant of Na,K-ATPase are more susceptible to hypoxic damage.

3.4. Mutant α1m-3CAβ1 (Cys454-458-459Ala) is presented with alterations in signaling function

Several mammalian species respond to hypoxia by release of cardiotonic steroids from adrenal glands into the circulation. These hormones interact with the Na,K-ATPase triggering activation of Src kinase [21]. We have tested the impact of Cys-to-Ala substitution on the receptor function of the Na,K-ATPase under normoxic and hypoxic conditions. Activity of Src kinase was used as a readout of signaling.
initiated upon the stimulation of Na,K-ATPase with low doses of cardiotonic steroid ouabain. Selection of optimal concentrations of ouabain was performed based on the previous studies in which dose-response of Src activity was measured as a function of ouabain concentrations in mouse fibroblast cells (SC-1 cell line). Maximal activation of Src in SC-1 cells was induced by treatment with 250 µM ouabain for 30 min under normoxic conditions [31]. We have thus chosen the same conditions of ouabain treatment for HEK293 cells expressing murine Na,K-ATPase α1 subunit. We also monitored the changes in Src activity in non-transfected HEK cells in the presence of 250 µM ouabain keeping in mind that this dose of ouabain is toxic as it blocks transport function of human Na,K-ATPase. 0.2% O₂ was used for acute (45 min) induction of signaling responses to hypoxia. Activation of Src kinase was assessed as the change in activating phosphorylation of Tyr416 and the activation of Erk1/2 kinases as a switch in Thr202/Thr204 phosphorylation. As expected, activating phosphorylation of Src kinase increased upon the stimulation with ouabain under normoxic conditions in α1m-wtβ1-carrying cells (Fig. 4A). Hypoxia alone was capable of activation of Src kinase to the same extent. This stimulatory response was suppressed if ouabain was present in deoxygenated cell culture medium (Fig. 4A). Similar response was observed in cells expressing α1m-1CAβ1 mutant. However, HEK293 cells expressing α1m-3CAβ1 and α1m-4CAβ1 enzymes were completely insensitive to the stimulation with ouabain at normoxic as well as at hypoxic conditions. HEK293 cells expressing endogenous human α1 subunit alone were not responding to 250 µM ouabain with modulation of Src activity, but retained hypoxia-sensitivity (Fig. 4A).

Erk1/2 activating phosphorylation was assessed in the same set of cell lysates to monitor signal transfer to the downstream elements of the Src-signaling cascade. In HEK293 cells carrying α1m-wtβ1 Na,K-ATPase the changes in activating phosphorylation of Erk1/2 mirrored that of Src (Fig. 4B). In these cells hypoxia induced phosphorylation of Erk was similar to that produced by exposure of cells to 250 µM ouabain under normoxic conditions, whereas a decrease in phosphorylation of Erk kinase was observed upon stimulation of hypoxic cells with ouabain (Fig. 4B). Substitution Cys444Ala was not affecting responses to ouabain under normoxic conditions but abolished phosphorylation of Erk kinase upon deoxygenation. Substitutions Cys454-458-459Ala made Erk1/2 less sensitive to both ouabain and low oxygen (Fig. 4B). Thus, cysteine residues within the NBD are critical in defining the signaling responses of the enzyme to ouabain and hypoxia.

3.5. Modeling predicts the critical importance of Cys 458, 459 within the NBD for interaction with kinase domain of Src

According to the reports of Li et al. [33] NBD of the Na,K-ATPase interacts with the kinase domain of Src kinase through the residues 410–429 of the α1 subunit. A short peptide NaKTide of identical sequence was effectively blocking interaction of Src kinase with Na,K-ATPase [33]. Cysteines 454, 458 and 459 within the NBD that turned to be essential for this interaction in our study (Fig. 4A) are localized out of the frame of NaKTide sequence (Fig. 5). We hence produced the model of the complex structure of kinase domain of Src kinase with the NBD of Na,K-ATPase in silico. The model was built based on the porcine Na,K-ATPase crystal structure (PDB code 3wgu) and human Src structure (PDB code 2arc). The NaKTide sequence was utilized for selection of structures of Src-NBD complexes.

Out of 40 models of the complex generated by docking, the model chosen for analysis showed maximal interaction of Src kinase with NaKTide without steric difficulties in reconstruction of full-size protein structures (Fig. 5A,B). In this model the interface of interaction was formed by the NBD residues 399-QS-400, 405-DKT-406, 466-RY-467, 488-NPNTAEPHR-497, 518-GK-519, 550-HLF-PDE-556 (numbering according to 3wgu). Cysteines 456 and 457 (458 and 459 in mouse sequence, highlighted in bold) formed an interface between the NBD of the Na,K-ATPase and the kinase domain of Src kinase. Cysteine 452 (454 in mouse sequence) was inward-facing and interacted with the residue 441 within NaKTide. Within the NaKTide sequence 408-SATWLALSRIAGLCNRAVFQ-427 the six residues were identified as interacting with Src (highlighted in bold). Within the Src sequence the interface between its kinase domain and the NBD of the ATPase exceeded that with NaKTide substantially and included the following amino acids: A165, 268-RLE; V314, 340-YMSK-343, 347-LKLG-350, 354-Q-355 the six residues were placed within the cavities proximal to these residues as discussed elsewhere [43]. S-glutathionylation of Cys 458 and Cys459 was simulated as we have described earlier [54]. Structure of S-glutathionylated protein was minimized in the Amber99 force field. Predictions for the interactions between the NBD of Na,K-ATPase and Src kinase were obtained with servers ClusPro, GrammX and ZDock. The resulting 30 structures were analyzed and 5 of them without steric difficulties in reconstruction of full-size protein structures were finally

![Fig. 3. Sensitivity of wild type and mutated murine Na,K-ATPase to hypoxia. A: Sensitivity of mouse wild type, wild type, α1m-1CAβ1, α1m-3CAβ1 and α1m-4CAβ1 isoforms to hypoxic exposure (24 h at 1% O₂). Data are means ± SD: Viability of non-transfected HEK293 cells and those transfected with wild type, α1m-1CA, α1m-3CA and α1m-4CA plasmids under hypoxic conditions. Cell viability was assessed as number of cells permeable for propidium iodide. Data are means of 3 experiments ± SD, * stands for p < 0.05 compared to normoxic controls.](image-url)
selected. QASDOM server was used for analysis of these models of NBD-Src kinase interaction interfaces. Interface of interaction included the following NBD residues: 392 – 416, 430 – 438, and 454 – 467. The best score was obtained for the model in which residues 397-ENQSGVSFDKT-407, 431-ENLPIL-436, 454-ELCC-457 of the Na,K-ATPase's NBD interact with the residues 272-KLGQGCFGEVW-282, 298-KP-299, 422-AK-423, 427 K, 467-VN-468 of the kinase domain of Src kinase (Fig. 5). In this model interface of interaction of S-glutathionylated NBD with Src kinase does not include NaKTide domain. The model predicts that glutathione bound to Cys 458 and 459 of NBD will prevent the interaction of kinase domain of Src with the NaKTide sequence within the NBD due to streric difficulties.

4. Discussion

Using mutagenesis approach we have revealed the importance of cysteine residues 244, 458 and 459 in redox- and hypoxia-sensitivity of the Na,K-ATPase as well as in its control over Src and Erk kinases. These regulatory mechanisms converge into the survival success under hypoxic conditions.

4.1. Localization of regulatory cysteines within the cytosolic domains of the catalytic α1 subunit

The possible role of cysteines within the catalytic α subunit for the Na,K-ATPase function was assessed in several studies. Substitution of all 23 cysteines in the α1 catalytic subunit by serine and alanine resulted in expression of unstable enzyme with a 2-fold reduced Km for Na+ and a 50% increased Km for K+ [28]. The enzyme containing cysteine-less α1 subunit was largely retained in the ER and Golgi apparatus and degraded 3 days after the transfection[28]. At the same time substitution of single cysteine residues by serine or alanine revealed particular importance of equivalent of Cys 244 in rat/rabbit/duck α1 subunit for folding of the α subunit and the enzyme stability in HeLa cells[60]. Substitution of analogues of Cysteines 454, 458 and 459 was associated with a reduction in maximal turnover rate to about 25% [60]. I n HEK293 cells we managed to express functional protein containing α1 subunit lacking Cys244 or/and Cys 458, 459 and 459 (Suppl. Fig. 2).

In our previous studies binding of glutathione to the Cys 244, 454, 458 and 459 was associated with the loss of Na,K-ATPase hydrolytic activity [54]. We then suggested that these residues may contribute to the enzyme's redox- and oxygen-sensitivity as sites of regulatory S-glutathionylation [54]. Further evidence that S-glutathionylation of
some cysteine residues (206, 454, 601 and 700) within the α1 subunit occurs before the enzyme is folded in the ER was provided recently [43]. These cysteines cannot be de-glutathionylated unless the protein is unfolded, and thus cannot participate in acute modification of the enzyme activity, but most likely defines the enzyme folding and stability.

In this study we have revealed the particular importance of Cys244 for making the enzyme sensitive to GSSG. Cysteine residues 454, 458 and 459 within the NBD are less essential for Na,K-ATPase redox sensitivity (Fig. 2). This observation was somewhat unexpected. Our previous findings indicated that inhibition of the enzyme by GSSG results from the outcompeting of ATP from its binding site upon S-glutathionylation [53,54]. Cys244 is localized within the actuator domain that does not contribute to the ATP binding site of the enzyme in E1 conformation [44]. In contrast to ADP, ATP binding to Na,K-ATPase generates a structural transition in the enzyme which causes transformation of the enzyme from the “E1-open” to “E1-closed” conformation with the following phosphorylation step [53]. Similar response was earlier described for GTP-induced conformation transitions in translational GTPases. Together with the earlier studies involving mutagenesis [59] and the observations on the sensitivity of the enzyme to oxidation [10] our present findings (Figs. 2 and 3) suggest that Cys244 joins the NBD in certain conformations as schematically shown in Fig. 6A.

The second unexpected finding of this study is the role of the cysteine residues within the NBD in the signaling mediated upon treatment of mouse Na,K-ATPase with ouabain.

4.2. Cysteines and the sensitivity of the Na,K-ATPase to ouabain

High salt diet [40], psychological stress [22], intensive physical exercise [3], and reduction in O2 availability [18] trigger release of cardiotonic steroids from the adrenal gland and hypothalamus into the circulation and initiation of multiple systemic physiological response [45,58]. Binding of ouabain to the regulatory binding site at the extracellular part of Na,K-ATPase α subunit stabilizes the enzyme in the E2-P conformation and initiates multiple events. Those include alteration of Src kinase activity, activation of inositol 3-phosphate receptors, induction of low frequency Ca2+ oscillations, and alteration in protein synthesis [56]. Src kinase is considered to be a part of the signaling complex [1,24,34,63]. The existence of distinct “signaling pool” of the Na,K-ATPase controlling the activity of Src kinase along with “ion pumping pool” was recently suggested [35]. Two binding sites for Src SH2 and kinase domains are localized within the actuator and NBD domains of the α subunit of the Na,K-ATPase respectively [2,33,34]. Src kinase remains inactive when its kinase domain is bound to the NBD. Strength of interaction between the SH2 domain of Src and actuator domain of the Na,K-ATPase exceeds the one between the NBD and Src kinase domain, suggesting that this interaction may be maintained upon dissociation of the Src kinase domain from its binding site within NBD in response to ouabain binding to the Na,K-ATPase [2]. Our data are in line with this hypothesis and imply that cysteines within the NBD are essential for the preservation of Src activation upon ouabain treatment (Fig. 4). Downstream events following the interaction of Na,K-ATPase with ouabain are defined by the environmental O2 availability (Fig. 4). Hypoxic exposure alone is sufficient for activation of Src kinase (Figs. 4, 6B, [29,31]). Exposure to ouabain under hypoxic conditions results in suppression of free radical production and inhibition of Src activity (Figs. 4, 6B, [31]). Replacement of cysteine residues within the NBD by alanine uncouples activity of Src from O2 availability or ouabain stimulation leaving the kinase permanently active (Figs. 4, 6C). Substitution of Cys244 with alanine does not affect ouabain- and hypoxia-driven regulation of Src activity (Fig. 4). These observations cannot be explained within the model of interaction between the kinase domain of Src kinase and NBD proposed earlier.
4.3. Cysteines, cell survival and responses of the Na,K-ATPase to hypoxia and 459 (Fig. 4).

According to Li et al. [33] Src binding site includes amino acids 410–429 of the α1 subunit. Our findings imply that Cys 454, 458 and 459 of NBDC is essential for the interaction with Src kinase and required for Src inhibition. This suggestion is further strengthened by the outcome of in silico modeling of binding of kinase domain of Src kinase to the NBD of the Na,K-ATPase (Fig. 5). According to this model Cys 454, 458 and 459 form the interaction interface between the Na,K-ATPase and Src kinase and binding of glutathione to them compromises the interaction (Fig. 5). Further evidence confirming the importance of cysteines of the NBD for the signaling function of the Na,K-ATPase comes from the analysis of the isoform-specificity of the receptor function of the enzyme. Whereas α1-containing isoforms are well-known for being involved in the regulation of Src activity, the enzyme formed by the α2 subunit was shown to be insensitive to the stimulation with low doses of ouabain [65]. From the three cysteines within the NBD only two are preserved in the α2 isoform of the catalytic subunit, namely the 454 and 459 whereas Cys 458 is not conserved. The α2 subunit also differs from the α1 in that it has one extra cysteine residue, Cys 236, in the actuator domain of the ATPase [10]. It would be tempting to assume that the presence of Cys 458 is critical for the signaling activity of the Na,K-ATPase. The enzyme formed by the α3 isoform of the catalytic subunit responds to the stimulation with low doses of ouabain with activation of Erk 1/2, but not Src kinases [39]. This interesting finding seems to contradict the “classical Xie model” describing Erk 1/2 activation as a downstream event following the Src activation upon ouabain binding to the enzyme [55]. In our study, uncoupling between the Src and Erk responses to ouabain and hypoxic stimulation was observed in cells lacking the three cysteines within the NBD (Fig. 6) suggesting the existence of an alternative Na,K-ATPase-driven, but Src-independent, signaling pathway involving Erk stimulation. Similar uncoupling between the Src and Erk activity has been reported earlier in cardiac myocytes [20,39,64] where Erk kinases could be activated via the IP3-dependent signaling cascade that was not necessarily related to the activation of Src by ouabain. Our data suggest that this uncoupling does not occur in cells expressing wild type mouse α1 isoform, but only in cells deprived of cysteines 244 and/or 454, 458 and 459 (Fig. 4).

We were able to confirm that exposure of HEK293 cells to low
oxygen results in suppression of the Na,K-ATPase hydrolytic activity (Fig. 1). Substitution of any of the cysteines we have chosen as candid-dates for regulatory cysteines to alanine interfered with the inhibitory effect of hypoxia observed in non-transfected cells as well as in cells expressing wild type mouse α1 protein (Fig. 3). Hypoxia induces an increase in the intracellular GSSG levels (Petrushanko 2012, Yakushev 2012). Data shown in Fig. 2 indicate that this inhibitory effect of GSSG on Na,K-ATPase is due to S-glutathionylation of Cys244. Increased hydrolytic activity of the Na,K-ATPase in mutants lacking Cys244 may be related to the ATP depletion and compensatory increase in ATP af-finity of the enzyme in these cells as has been observed earlier in re-sponse to GSH-depletion [50]. Further work is required for in-depth characterization of the mechanisms of the inhibitory action of hypoxia on the Na,K-ATPase that may be rather complex as recently reviewed [11].

We have revealed that hypoxic exposure of HEK293 cells as well as of HEK 293 transfected with α1wt or α1m-1CAβ1 resulted in acti-vation of Src kinase. Signaling responses to ouabain in α1wt and α1m-1CAβ1-transfected HEK293 were inverted under hypoxic condi-tions (Fig. 4). Replacement of Cys 454, 458 and 459 by alanines render-ed Src kinase permanently active and insensitive to ouabain and hypoxia (Fig. 4A). Data shown in Fig. 5D suggest that permanent ac-tivation of Src in hypoxic cells hypoxia is due to glutathionylation of Cys458 and Cys459. Similar to Src, Erk kinase becomes independent of O2 availability and ouabain in the absence of cysteines within the NBD domain (Fig. 4). Preconditioning triggered by brief “pulses” of treatment of isolated perfused heart or brain slices with cardiotonic steroids (ouabain, digitoxin, nerifolin) could protect the tissue from damage caused by ischemia-reperfusion [20,4,49,5]. This protective action of ouabain in mouse heart relied on the of IP3 kinase-1A, and was in-dependent of activation of Src kinases [20]. Digitoxin was identified as a potent suppressor of autophagy-related apoptosis or necrosis (autosis) of hippocampal neurons in a rat model of cerebral hypoxia-ischemia [37]. Signaling pathways causing cytoprotective effects of cardiotonic steroids are currently unknown. They are most likely associated with the changes in free radical production within the mitochondria under hypoxic conditions that in turn may be intimately linked to the calcium filling of the ER and ER stress [1]. It is unclear whether activation of Src kinases under hypoxic conditions as well as in the presence of ouabain is caused or by themselves may cause the changes in free rad-ical production. Src kinases were earlier on shown to respond with activation to the free radical burst or exposure to peroxynitrite [25,30,42]. Free radical burst and the related increase in peroxynitrite production and tyrosine nitration were demonstrated during the first 10–30 min of hypoxia [26,66]. At the same time, hypoxia results in activation of Src kinase as it was shown earlier [29,31] and in this study (Fig. 4). Bringing these two observations together the model suggests that activation of Src is caused by S-glutathionylation of the cysteines within the NBD domain and the following de-attachment of Src kinase from the Na,K-ATPase. Treatment of hypoxic HEK293 cells expressing murine α1 subunit with 250 μM of ouabain resulted in inactivation of Src and promotion of its interaction with NBD of the Na,K-ATPase (Fig. 6). Reported earlier on, inhibition of Src caused by ouabain in SC-1 cells under hypoxic conditions was associated with a decrease in pro-duction of reactive oxygen species and prevention of oxidative stress [31].

Taken together assessment of signaling responses of Na,K-ATPase (Fig. 4) and the results of modeling (Fig. 5) indicate pivotal importance of Cys 454, 458 and 459 within the NBD in interaction between the Na,K-ATPase and kinase domain of Src. Replacement of these cysteines with alanine or their S-glutathionylation prevents formation of the “inhibitory” complex between the Src kinase and the Na,K-ATPase. Src remains permanently active and insensitive to hypoxic challenge or stimulation with ouabain. Interestingly, treatment of the enzyme in cell lysates with access of GSSG or overloading of cells with it causes a different pattern of responses in which Cys 244 is of foremost importance in control of the enzyme’s hydrolytic function (Fig. 2). We may hence only speculate about the possible interplay between both processes in hypoxic organism. Hypoxia triggers free radical burst o-riginating from the mitochondria [26]. Along with that deoxygenation causes release of cardiotonic steroids from the kidney adrenal glands and hypothalamus into the circulation. Endogenous ouabain is pro-duced from cholesterol in the mitochondria, the organelles involved in ATP production [36,62]. This response may be adaptive and convey protection against ischemia-reperfusion injury (see above). Fur-thermore, suppression of the Src activity triggered by ouabain under hy-poxic conditions should suppress proliferation under hypoxic condi-tions and preserve the cells from unnecessary high ATP consumption, the effect that powers the use of cardiotonic steroids as potential anti-cancer therapy [56]. Our findings indicate that this adaptive response may be lost in the absence of the Cys 244, 454, 458 and 459 of which Cys 458, 459 and 244 are likely to be of utmost importance (Figs. 4 and 6). These cysteine residues within the catalytic α subunit are most likely involved in control of ATP expenditure under hypoxic conditions by controlling the hydrolytic activity of Na,K-ATPase and its signaling function that extends much beyond the ATP consumption by the pump alone as stated in the “channel arrest theory” [12,19,27,47].

5. Conclusions

The results obtained emphasize the important role of regulatory cysteines located in the AD and NBD of the Na,K-ATPase in the ability of enzyme to respond to hypoxia tuning its hydrolytic activity and signaling function to the changes in environmental oxygen availability. For the first time the critical role of Cys244 was shown in inhibition of enzyme by regulatory S-glutathionylation. We also revealed the impor-tance of Cys458 and Cys459 in the interaction between the Na,K-ATPase and Src kinase. This interaction is a core event in receptor function of the enzyme during exposure to cardiotonic steroids and hypoxia.

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Appendix A. Supplementary material

Supplementary data associated with this article can be found in the online version at http://dx.doi.org/10.1016/j.redox.2017.05.021.

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