Prospects & Overviews

Revisiting Kadenbach: Electron flux rate through cytochrome c-oxidase determines the ATP-inhibitory effect and subsequent production of ROS

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Mitochondrial respiration is the predominant source of ATP. Excessive rates of electron transport cause a higher production of harmful reactive oxygen species (ROS). There are two regulatory mechanisms known. The first, according to Mitchel, is dependent on the mitochondrial membrane potential that drives ATP synthase for ATP production, and the second, the Kadenbach mechanism, is focused on the binding of ATP to Cytochrome c Oxidase (CytOx) at high ATP/ADP ratios, which results in an allosteric conformational change to CytOx, causing inhibition. In times of stress, ATP-dependent inhibition is switched off and the activity of CytOx is exclusively determined by the membrane potential, leading to an increase in ROS production. The second mechanism for respiratory control depends on the quantity of electron transport to the Heme aa3 of CytOx. When ATP is bound to CytOx the enzyme is inhibited, and ROS formation is decreased, although the mitochondrial membrane potential is increased.

Keywords:
- allosteric inhibition; cytochrome c oxidase; enzyme kinetics; ischaemic preconditioning; phosphodiesterase inhibitors; reactive oxygen species

Introduction

Mitochondrial respiration is the most common mechanism for ATP production and energy supply. Energy demand varies and depends on different forms of activities in cells, tissues and organisms. Early in earth’s history, there were Prokaryotes that were dependent on their own metabolism and on nutrients that were provided by their environment. Approximately 250 million years ago, after dramatic changes occurred in earth’s atmosphere, its composition changed to 21% oxygen [1]. The altered conditions led to a change in the production of cellular energy from anaerobic glycolysis to respiration because a huge energy supply was required for the survival of Eukaryotes [2]. The question of adequate regulation of respiration appeared. As is known today, regulation of respiration is absolutely essential to minimise potentially harmful by-products, including reactive oxygen species (ROS). The factors that adapt respiration to physiological strain remain poorly understood (Box 1).

Polarographic assay of CytOx activity enables detection of an ATP-dependent inhibitory effect

Respiratory steady states have already been defined by Chance and Williams [6] according to a protocol for oxygraphic experiments with isolated mitochondria corresponding to the activities of all 5 of the mitochondrial multienzyme complexes

Abbreviations:
- CytOx, cytochrome c oxidase (E.C. 1.9.3.1.); H, Hill slope coefficient (according to GraphPad Prism software); PEP, phosphoenolpyruvic acid; PK, pyruvate kinase (E.C. 1.7.1.40); ROS, reactive oxygen species; TN, turnover number.

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of the ETC. Ferguson-Miller et al. [7] reported the polarographic measuring procedure in a measuring cell, which operates according to the principle of the Clark electrode, which is based on selective electron transfer by an electron donor (ascorbic acid 18 mM) and an electron transmitter (cytochrome c in increasing concentrations) to the mitochondrial respiratory chain complex IV (CytOx). Subsequently, Kadenbach and co-workers used this system to perform measurements either in the presence of 5 mM ADP or 5 mM ATP. For measurements in the presence of ATP, an ATP regenerating system (10 mM Phosphoenolpyruvate, 2 U/mL pyruvate kinase, 5 mM MgSO₄) was also used to maintain the ATP concentrations high enough and to demonstrate the effect of inhibited CytOx.

Studies by Arnold and Kadenbach [8] described the influence of intramitochondrial ATP/ADP ratios with increasing amounts of cytochrome c in the liposomally reconstituted enzyme. An increased ATP to ADP ratio resulted clearly in sigmoidal enzyme kinetic curves (Hₑ from 1.09 to 1.97) at increasing cytochrome c concentrations (from 0.25 to 60 μM). However, at high concentrations of cytochrome c, the enzyme kinetics became hyperbolic. In these experiments, CytOx kinetics was measured as ‘Turn over number’ (TN [S⁻¹]) [9, 10]. CytOx was reconstituted in proteoliposomes at a concentration of 50 nM with varying ratios of ATP/ADP inside the vesicles. Therefore, the aa₃ content inside the proteoliposomes was constant, but addition of increasing amounts of cytochrome c resulted in a steady increase of electron transmissions to aa₃ and in higher TN’s at the end. According to our hypothesis, the shift from sigmoidal to hyperbolic kinetics described in this report is attributed exclusively to the electron transmission rate to CytOx, whereas the enzymatic consumption of oxygen itself depends on the uptake of electrons by aa₃ when increasing amounts of cytochrome c are transferred to dioxygen for reduction [9, 10].

Acceptance of this type of CytOx activity measurements has always been controversial. First, criticisms were based on the fact that original measurements from Arnold and Kadenbach [8], which were performed using a reconstituted enzyme, were not directly comparable to measurements with the isolated enzyme, or measurements with mitochondria or tissues. Second, the common use of N,N,N',N'-tetramethyl-p-phenylenediamine (TMPD) in enzymatic kinetic measurements hides the electron transfer rate dependency because electrons are transferred not only to cytochrome c but also directly to the enzyme [11] and even bypass the cytochrome c binding site [12]. Third, the use of detergents, which is thought to have intermediary effects on enzymatic kinetic measurements for permeation of cytochrome c through the outer mitochondrial membrane, is rather deleterious. Detergents destroy the mitochondrial membrane’s architecture and ‘create’ a mixture of isolated enzymes, mitochondrial and tissue fragments as well. Different forms of agglutinates were observed by electron microscopy. The activity of membrane bound enzymes varies in a wide range. The activity of the purified enzyme is partially inhibited by
Box 2

Current view of Kadenbach’s theory as an extension of Mitchell’s theory

The molecular mechanisms are currently not fully understood. With regard to the present concepts see Fig. 2A–D.

1. As is generally known, electrons are transferred through the ETC (from complex I to IV). Additionally, complexes I, III and IV act as proton pumps, translocating hydrogen ions (protons) across the inner mitochondrial membrane and creating a mitochondrial membrane potential. This $\Delta \psi \text{m}$ drives complex V (ATP-synthase) toward production of ATP. As reported by Kaim and Dimroth, an optimal range between $80 \text{mV} < \Delta \psi \text{m} < 120 \text{mV}$ guarantees sufficient synthesis of ATP, resulting in the transfer of 4 electrons to dioxygen (reduction) and oxidation of hydrogen to water without excessive production of ROS. At $\Delta \psi \text{m} > > 120 \text{mV}$, large amounts of ROS are produced [5, 63]. In our measurements using isolated myocardial mitochondria, we have found greater increased values of $\Delta \psi \text{m}$, thus we assume a permanent release of ROS under such experimental settings in vitro.

2. Under physiological conditions, the cell contains extremely high quantities of ATP. In the so called ‘relaxed’ state, ATP binds to subunit IV of the phosphorylated CytOx and induces an allosteric conformational change that results in sigmoidal enzyme kinetics. Therefore, CytOx is ‘ATP-dependent allosterically inhibited’ and an increased production of ROS during respiration is avoided. Although the membrane potential is sustained by the ATP demand, in this state (within a stable range of $\Delta \psi \text{m}$), the enzymatic activity of CytOx after ATP binding is regulated exclusively by the mitochondrial ATP/ADP ratio.

3. In the case of cellular stress, allosteric inhibition of the enzyme by ATP is switched off. Although the maximum rate of ATP synthase is known to be beyond $100–120 \text{mV}$ [63], this $\Delta \psi \text{m}$ increases because of the higher CytOx activity. Because CytOx is the rate-limiting step in the ETC, its activity increases for higher ATP production [64]. Electron transmission from complexes I to IV is accelerated at closer distances between the complexes because of mitochondrial membrane shifts [65], resulting in maximum production of water with CytOx. Both prokaryotic and eukaryotic ATP synthase complexes have the same capacity with respect to their maximum rates [71]. A turnover number for ATP synthesis of $270 \pm 40/\text{s}$ was determined in the presence of 5% active F0F1 complexes. In the case of hyperpolarization of $\Delta \psi \text{m}$, the potential drives ATP synthase beyond its capacity, resulting in the maximum production of ATP, but exceeds effective ‘oxygen utilization’. The $\Delta \psi \text{m}$ exceeds the normal range, and ROS are formed in high concentrations because of the maximum synthesis rate of ATP synthase [63]. At this stage, CytOx activity is determined by $\Delta \psi \text{m}$, and no longer by the ATP/ADP ratio. In parallel, the concentration of ROS increases, resulting in harmful effects to the cell.

4. Kadenbach’s Hypothesis states that the regulation of the membrane potential and ROS formation in mitochondria are determined by the ATP-induced allosteric inhibition of CytOx, and represents a second mechanism for respiratory control. Under relaxed conditions, feedback inhibition of CytOx by ATP maintains the membrane potential at low values. Stress factors increase the cytosolic and/or mitochondrial [Ca$^{2+}$], which activates calcium-dependent protein phosphatases and dephosphorylates CytOx. Without allosteric inhibition of CytOx by ATP, the membrane potential increases with a consequent increase of ROS formation. Question regarding initial mechanisms have yet to be answered. There is a competition between ADP and ATP for the binding sites on the enzyme [66]. With reconstituted enzyme, the kinetics were influenced by extraliposomal (cytosolic) ATP and ADP. The Km for cytochrome c was five times higher in the presence of extraliposomal ATP than with ADP. These differences of Km values were abolished after preincubation of the enzyme with a monoclonal antibody to subunit IV. The data demonstrate the regulation of cytochrome c oxidase activity by the cytosolic ATP/ADP ratio, in addition to regulation by the matrix ATP/ADP ratio. Cyclic AMP activation of mitochondrial PKA, which is generated by the carbon dioxide/bicarbonate-regulated soluble adenyl cyclase [67] is found to induce phosphorylation of CytOx [68] and to influence the enzymatic activity. Finally, the redox-dependent transfer of protons to the binuclear centre through the D-channel and the K-channel, where the latter is redox-independent [69] remains to be clarified. Interacting effects of $\Delta H^+$ and $\Delta \psi \text{m}$ on $\Delta \text{pH}$, as a controlling step in the ETC activity have already been discussed [70]. Nevertheless, our data suggest that there is no allosteric inhibition of CytOx when electron transfer is increased. Permanent stress and elevated ROS levels can result in cell apoptosis and the generation of multiple degenerative diseases.

Triton X-100 and dramatically enhanced by Tween 80 or phospholipids [13].

The latter two factors especially were considered for a long time as interfering factors for efficient kinetic measurements of the enzyme. Effective measurements are achieved only under very stringent conditions. We performed very simple experiments (supplemental data in [14]) using bovine heart tissue homogenate and isolated bovine heart mitochondria (Fig. 1A and B), as well as rat heart tissue homogenate and isolated rat heart mitochondria (Fig. 1A–D). In all cases, measurement conditions were standardised as mentioned previously [14], and reproducible results were achieved.
Interestingly, measurements using homogenates in both cases consistently demonstrated significant allosteric ATP-dependent enzyme inhibition, in contrast to results with isolated mitochondria. The term 'ATP-dependent enzyme inhibition' is primarily observed at low, rather physiological intramembrane concentrations of Cytochrome c [15, 16], and subsequent to the performed kinetic experiments. In these experimental settings, spectrophotometric analysis revealed a difference in aa3 content at the same protein ratio of 1–2.25. When H > 1 as shown in Table 1, the kinetics of the enzyme are allosteric as is apparent in Fig. 1A–D. When H < 1, the curve is sigmoidal due to positive cooperativity, as in the presence of ATP and an ATP-regenerating system. We observed sigmoidal curves during tissue homogenate measurements of both rat heart and bovine heart. (Table 1, Fig. 1A–D). Consequently, polarographic measurements performed with different concentrations of freshly isolated mitochondria showed altered enzyme kinetics with dilution series of mitochondria (Ramzan et al., unpublished results). Variations of involved components (ascorbate, cytochrome c) confirmed these results. Therefore, we conclude that the 'Kadenbach effect' is triggered by two main components: (i) the extremely high amount of ATP that is present in intracellular high-energy phosphates, and (ii) the number of electrons transmitted from cytochrome c to aa3 of CytOx.

Table 1. Using an allosteric sigmoidal programme in GraphPad Prism software, Hill-slope (H) calculations were performed using the CytOx kinetics shown in Fig. 1A–D

| Samples                               | ADP (H) | ATP (H) |
|---------------------------------------|---------|---------|
| Bovine heart tissue homogenate (n = 4) | 0.8899  | ~1.103  |
| Bovine heart mitochondria (n = 4)     | 0.9802  | 0.8115  |
| Rat heart tissue homogenate (n = 3)   | 0.5566  | 1.152   |
| Rat heart mitochondria (n = 3)        | 0.8868  | 0.8062  |

When H > 1 as shown in Table 1, the kinetics of the enzyme are allosteric as is apparent in Fig. 1A–D. When H < 1, the curve is hyperbolic in accord with the Michaelis-Menten equation.
molecular cause of cardiac injury after ischaemia must focus on mitochondria [18, 23, 24] and their production of ROS.

Kadenbach’s theory provides an explanation for the regulation of ROS production in mitochondria. ROS are considered the main cause of development of degenerative diseases [25], for damaging the heart after ischaemia [26], and in development and progression of heart failure (Box 3) [27, 28–32].

The production of ROS is dependent on the regulation of the mitochondrial membrane potential (ΔΨm). The theory of Kadenbach describes a mechanism that maintains the ΔΨm at low values under normal conditions, thus preventing excessive formation of ROS. This mechanism is switched off under conditions of stress and excessive work to maximise the rate of ATP synthesis and is accompanied by a decreased efficiency [56]. There is a physiological balance between oxidative and reductive processes in biological systems, and the constant relationship between both is maintained by an ‘antioxidative capacity (AOC)’. A predominance of reactive oxygen intermediates is referred to as ‘oxidative stress’. ROS are scavenged by corresponding enzyme systems that are classified as either enzymatic or non-enzymatic [57, 58]. This classification has clinical value because the regulation of redox reactions is important for protecting the heart from coronary disease [59, 60]. These observations conform to the work of Prosser et al. [61] who found that production of ROS is induced when cardiac cells are physiologically strained. Additionally, the induction of protein neogenesis and protein assembly of mitochondrial proteins has also been assumed (Box 3) [62].

**Signaling factors and CytOx interactions**

The basis of Kadenbach’s theory is the proposal of a ‘second mechanism of respiratory control’ [50] that regulates the rate of respiration by the ATP/ADP ratio, extending the ‘first mechanism of respiratory control’, whereas excessively high values of ΔΨm limit the respiration rate. Arnold and Kadenbach have shown that high intramitochondrial ATP/ADP ratios convert the hyperbolic kinetics of ascorbate-dependent respiration of isolated CytOx to an inhibited allosteric kinetic status (Hill coefficient >1), which is independent of the ΔΨm and is based on the binding of ATP or ADP to the matrix domain of CytOx subunit IV [72]. Ogbi et al. [73] observed that phosphorylation of CytOx subunit IV by protein kinase Cα produced an increase in the CytOx activity. They demonstrated that 4-phorbol esters activate translocation of PKCs into mitochondria, which were immunoprecipitated with the CytOx subunit IV. The role of other subunits in this mechanism remains to be clarified. The subunits IV, VIa, VIb, VIIa and VIII are nuclear-encoded subunits of CytOx and are expressed in 2 or 3 tissue- or development-specific isoforms [74–76]. The catalytic centre of the enzyme is located in three subunits that are encoded in the mitochondrial genome (haeme a and haeme a3/CuB in subunit I CuA in subunit II, see Fig. 2C and D). In addition to the catalytic subunits, the mammalian enzyme also contains 10 subunits that are encoded by the nuclear genome, causing complex regulation of the enzymatic activity [52]. Whether phosphorylation of subunits is involved, is still an open question. Western blot analysis of isolated CytOx (isolated under different conditions), using antibodies against phosphoserine and phosphothreonine, identified multiple phosphorylation sites on subunits I, II, III, IV, VIa, VIb, Vlc, VIIa, VIIb and VIIc [9, 10]. Allosteric inhibition of ATP is reversed by binding of thyroid hormone 3,5-diido-L-thyronine, to CytOx subunit Va [77]. Surprisingly, Lee et al. [78] observed that AMP-dependent phosphorylated enzyme exhibited only allosteric ATP inhibition, which is abolished by Ca2⁺-activated
Figure 2. Schematic representation of the modified molecular structure of CytOx subunits, originally from Herrmann et al. [123]. The mitochondrial-encoded SU I, II and III have the central stage, whereas the nuclear-encoded SU surround the central column. The blue arrow represents the binding of oxygen to the transmembrane helices of SU I and II [124]. Cytochrome c (molecule on the left, Hoffmeister K, Wikimedia commons) transfers electrons to CytOx (grey arrow ‘e−’). It is proposed that during enzyme turnover the enzyme cycles between two conformers, one with a substrate binding site on subunit II, and the other along the interface of subunits II, IV and VIb. Structural analyses suggest that Glu112, Glu113, Glu114 and Asp125 of subunit IV, and Glu40, Glu54, Glu78, Asp35, Asp49, Asp73 and Asp74 of subunit VIb are residues that could possibly be involved [125]. Cytochrome c binding affects the conformation of cytochrome a within CytOx [126]. A: Proposed model representing the influence of ADP or ATP binding to SU IV and SU VIII on the enzymatic activity of CytOx. Ten binding sites for adenine nucleotides are known. At seven sites, ADP and ATP are exchanged [127]. One binding site for ATP or ADP, located at the matrix-oriented domain of the heart-type subunit VIaH, increases the H+/e− stoichiometry of the enzyme in heart or skeletal muscle from 0.5 to 1.0 when bound ATP is exchanged by ADP. Two further binding sites for ATP or ADP are located at the cytosolic and the matrix domain of subunit IV. Although the additional binding site on SU VIa has been confirmed by Taanman et al. [128] (not shown), most binding sites were found on SU IV and VIII using radioactive ATP analogues, suggesting that these two nuclear-coded polypeptides may play a regulatory role [129, 130]. Especially, SU IV is essential for the assembly and respiratory function of the complete enzyme complex [131]. Because of the negative charges associated with ATP (fourfold), and the dipole moment of cytochrome c [132, 133], the holoenzyme creates an electrostatic field (negative sign on the cycle) that finally regulates the internal electron-transfer reactions by its electric field strength [134]. This explains how CytOx acts like an ‘electro-catalyst for oxygen reduction’ [135]. Furthermore, Craig et al. [136, 137] and Lin et al. [138] found that ATP binding to cytochrome c diminishes electron flow in the mitochondrial respiratory pathway and respiration is shut down. B: In the case of the exchange of ADP to ATP on the seven nucleotide binding sites the electrostatic field becomes weaker because of less negative charge with ADP. Subsequently, electron transfer from cytochrome c to SU II becomes accelerated. C: Modified model for subunit order inside the CytOx molecule according Tsukihara et al. [139, 140] and shows again the proposed mechanism of ATP binding to SU IV and VIII. The subunits of CytOx in the molecule centre are shown with blue (SU I), pink (SU II) and dark grey cycles (SU III). Roman numbers represents the helices. Blue dotted lines mark the entry of Helix I/II/III to Oxygen pathway 1 and the entry of Helix IV/V to Oxygen pathway 2. The binding of ATP (small grey cycles with white minus signs) at seven positions to SU IV and VIII results in a higher negative charge for the molecular dipole. The more negative ‘cloud’ induces tilting and bending of the molecule, and the binding of cytochrome c (black dotted line) is influenced, resulting in alterations of the subunit positioning (here helices XI, XII, I and II) together with a reduction in the distance between haeme a and haeme a3. The influence of an electric potential field and the effect of ionic strength on the reaction rate of cytochrome c have been described by Koppenol et al. [141]. D: The same molecular model features the situation after binding of ADP to all the binding sites of SU IV and VIII. A less negatively charged ‘cloud’ (left side) widens the distances between Helices XI, XII, I and II and finally induces a ‘more open’ angle between haeme a and haeme a3 for acceleration of electron transfer and increased Dioxygen turnover. However, the question of a pH-dependent polarity change at the binuclear centre [142] remains unanswered, although the proton K-pathway is known to become sufficiently flexible for internal water molecules to alternately occupy upper and lower parts of the oxygen pathways, which are associated with conserved Thr-359 and Lys-362 residues. Subsequent intramolecular ‘constrictions’ [143] could support the already known effect of dielectric relaxation of CytOx [144].
ATP-dependent allosteric inhibition, allostery and phosphorylation sites of CytOx: Data remain controversial

Iksoo Lee was the first to demonstrate in her thesis an obvious correlation between the phosphorylation of cytchrome c oxidase by a cAMP-dependent protein kinase A and ATP-dependent allosteric enzyme inhibition [9, 10]. By comparing consensus sequences, she suggested that this effect is triggered by phosphorylation of serine 441 in subunit IV. In contrast, Hüttemann and co-workers [89] identified cAMP-mediated inhibition of the enzyme, probably due to phosphorylation of tyrosine 304 on subunit I. An illustration from this original work; however, clearly shows that this phosphorylation is responsible for the rightward shift of the kinetics of the enzyme activity leading to the sigmoidal allosterically inhibited state but not to the theoretically expected transition of the enzyme kinetics to a hyperbolic state (Michaelis Menten kinetics 1st order). Another important observation was reported by Arnold and co-workers [90]. By a gradual reduction of oxygen, it was shown that transcription of CytOx subunit IV-2 was induced specifically in astrocytes. Increased transcription of isofrom IV-2 caused an obvious switching off of the allosteric inhibition mechanism for CytOx in the presence of high concentrations of ATP. It was concluded that the presence of this isoform removes allosteric inhibition of the enzyme due to a reduced responsiveness to the allosteric regulator ‘ATP’. Therefore, an oxygen sensor function has been assigned to CytOx. Of course, different phosphorylation sites on the enzyme were examined in terms of their functional relevance [68], however, the identification and importance of the individual phosphorylation site responsible for the ATP-dependent allosteric inhibition of CytOx remained in doubt.

Hüttemann and co-workers [89] claimed that the allosteric ATP-dependent inhibition of CytOx from bovine liver is related to the cAMP-dependent phosphorylation of tyrosine 304 on the cytosolic side of the subunit I. Miyazaki et al. [91] demonstrated phosphorylation of subunit II of CytOx by a non-receptor tyrosine kinase c-Src in osteoblasts and found a positive correlation between CytOx activity and c-Src kinase activity, although the amino acids that were phosphorylated remained obscure. A specific non-receptor tyrosine phosphatase, SHP-2, was detected by Salvi et al. [92] in mitochondria. Steenart and Shore [93] performed in vitro phosphorylation of CytOx subunit IV with [γ-32P] ATP, but did not identify the phosphorylated amino acid. The signaling pathways leading to phosphorylation and modification of CytOx activity are still largely unknown. Hüttemann and co-workers [89] have shown that the phosphorylation of Y304 in the CytOx subunit I is performed through G-protein-dependent receptors and that tyrosine phosphorylation of subunit IV is probably induced via the PI3 K (phosphatidylinositol 3 – kinase)/Akt (protein kinase B) pathway. Bijur and Jope [94] demonstrated phosphorylation of subunit II of CytOx by a non-receptor tyrosine kinase c-Src in osteoblasts and found a positive correlation between CytOx activity and c-Src kinase activity, although the amino acids that were phosphophorylated remained obscure. A specific non-receptor tyrosine phosphatase, SHP-2, was detected by Salvi et al. [92] in mitochondria. Steenart and Shore [93] performed in vitro phosphorylation of CytOx subunit IV with [γ-32P] ATP, but did not identify the phosphorylated amino acid. The signaling pathways leading to phosphorylation and modification of CytOx activity are still largely unknown. Hüttemann and co-workers [89] have shown that the phosphorylation of Y304 in the CytOx subunit I is performed through G-protein-dependent receptors and that tyrosine phosphorylation of subunit IV is probably induced via the PI3 K (phosphatidylinositol 3 – kinase)/Akt (protein kinase B) pathway. Bijur and Jope [94] demonstrated phosphorylation of Akt after activation of PI3 K by IGF-1 (insulin-like growth factor) in cell cultures (SH-SY5Y, HEK293) and further demonstrated that the phosphorylated Akt is translocated into mitochondria where it phosphorylates the β subunit of ATP synthase, glycojen synthase kinase-3β and other unknown proteins. The Manfredi group identified another important pathway for cAMP action concerning regulation of oxidative phosphorylation [67]. They proposed an intramitochondrial CO2-HCO3- → sAC-cAMP-PKA regulatory pathway for oxidative phosphorylation. The latest findings of Hess et al. indicate a ’CO2⁻ HCO3⁻ → sAC-cAMP- signalosome’ that is responsible for PKA activation and phosphorylation of subunit Va at positions T65 and S 43 of CytOx in Saccharomyces cerevisiae under normoxic conditions [95]. These phosphorylations modulate the allosteric regulation of CytOx by ATP and the authors showed that the normoxic subunit Va is a homologue of human subunit IV-1 (isoform), but the same experiments in human systems have yet to be performed. Acín-Perez et al. [96] demonstrated that residue S56 in mammalian CytOx subunit IV-1 is coupled with the prevention of allosteric inhibition of CytOx by ATP. In addition to discussions concerning
phosphorylated residues of CytOx [97] these data demonstrate the allosteric inhibition of CytOx by ATP and confirms part of Kadenbach’s theory. We have already shown a relationship between the ATP-dependent inhibition of CytOx and decreased ROS production [98]. Finally, the question remains whether all the ATP-dependent inhibitory effect of CytOx is always associated with allostery and for additional factors causing allostery.

Yaniv et al. [99] found that cAMP/PKA signaling is dependent on Calcium regulation. Effects on mitochondrial metabolism are due to the activation of soluble mitochondrial Adenylyl Cyclase by bicarbonate and calcium [100]. However, conflicting data were also published by the Balaban group. They observed a stimulation of oxidative phosphorylation by calcium without an influence by cAMP and PKA activity [101]. The pH dependency of bicarbonate-regulated soluble Adenylyl Cyclase [102] remains to be clarified in the context of the inhibitory effect of ATP on CytOx. Finally, Aciñ-Perez et al. [103] described a Phosphodiesterase 2 A that is localized in mitochondria and is involved in the regulation of respiration. This type of PDE2A is located in the matrix. Concerning different signaling chains for protein phosphorylations [104] and multiple phosphorylation sites of CytOx [105, 106], and the ‘so far known’ compartmentation of cyclic nucleotide signaling [107] on the other hand, we have to address the question whether all the different cAC actions [108] are maintained by a network of different PDE’s in the mitochondria or in the intramembranous space [109].

**Phosphodiesterase inhibitors as true regulators?**

regarding the data from the Manfredi group, Lee and co-workers studied signaling pathways targeting mitochondria and examined phosphorylation of CytOx subunits by the cAMP-dependent pathway. Using phospho-antibodies against phospho-tyrosine, they detected phosphorylated cow liver CytOx subunit I in the presence of theophylline, a phosphodiesterase inhibitor (PDE inhibitor) that induces high levels of cAMP. This type of phosphorylation of Tyr304 in CytOx decreased V(max) and increased K(m) for cytochrome c. It shifted the reaction kinetics from hyperbolic to sigmoidal as CytOx is fully or strongly inhibited up to 10 mM concentrations of cytochrome c [89]. Phosphodiesterase inhibitors (PDE) are known from their use in therapy of cardiovascular diseases, e.g. treatment of cardiac insufficiency. A wide spectrum of pharmaceuticals display their actions directly or indirectly on the status of mitochondrial bioenergetics. Surprisingly, our research group observed that the drugs Milrinone (PDE III inhibitor; 2-methyl-6-oxo-1,6-dihydro-3,4'-bipyridine-5-carbonitrile) and Euphylong (Theophylline; 1,3-Dimethylxanthin) had an opposite effect on CytOx kinetics (Fig. 3A–D). Allosteric inhibition was intensified by Milrinone, whereas Theophylline reversed this inhibition completely. These beneficial effects of Theophylline on ischaemic tissues act in a dose-dependent manner [110]. Milrinone treatment in cases of severe cardiac failure appears in a new spotlight [111] because myocardial dysfunction after ischaemia/reperfusion [35, 112] could be prevented by administration of Milrinone [113]. PDE networks appear confusing. Inhibitors of PDE, which cause increased concentrations of cyclic nucleotides, are expressed in multiple tissue-specific isoforms [114]. Until recently, 21 human PDE genes had been identified with...
Conclusions and outlook

The inhibition of CytOx by ATP presents a ‘second mechanism of respiratory control’ [50], which regulates the respiration rate by the ATP/ADP ratio, supporting the ‘first mechanism of respiratory control’, whereas the respiratory rate is limited at high ΔΨm values. When ATP is bound to CytOx, the enzyme is inhibited, and there is subsequently low formation of ROS, although the mitochondrial membrane potential is increased. The initial data from the Kadenbach group were confirmed, whereas the sigmoidal kinetics of the enzyme at high intramitochondrial ATP/ADP ratios with ADP bound to the CytOx subunit IV and replaced by ATP remains to be clarified [72]. Allosteric inhibition by ATP indicates a blockage of CytOx enzymatic activity at high concentrations of ATP, which is likely removed by dephosphorylation of CytOx as a result of activation of a calcium-dependent protein phosphatase [79]. However, the different phosphorylation steps of CytOx subunits are not completely understood. The allosteric inhibition of CytOx by ATP represents a control circuit at low ΔΨm values, which maintains the mitochondrial membrane potential within a physiological range. It has been confirmed that ATP binding to CytOx diminishes electron flow in the ETC [116]. In experiments with 8-Azido-ATP-modified CytOx and with Cytochrome c, modulation of electron transfer from Cytochrome c to CytOx by interacting with the enzyme and allosterically altering the docking was confirmed. However, if binding of ATP affects primary Cytochrome c or CytOx or both, reduced electron transfer remains open [117–119]. However, in fact, the docking scenario of Cytochrome c to CytOx under the influence of ATP is changed. Whether the ATP-cytochrome c adducts have a different binding site or a different docking conformation remains to be demonstrated. It is worth noting that the influence of Cardiolipin is not negligible. Tuominen et al. [120] found that ATP induction of conformation alterations was dependent on binding of lipid to Cytochrome c via an Arg93-containing binding site. Cytochrome c bound to Cardiolipin and ATP has a high level of Peroxidase activity that favours protein structures with an open haeme pocket [121] (see Fig. 2A and B). Therefore, we suggest both an electron scavenging effect and a modification of subunit I containing the two haeme centres on CytOx by ATP binding (see Fig. 2A and B). Haeme a acts as an ‘opened or closed baseball glove’ catching as an electron input device. Haeme a3 acts as part of the binuclear centre and site of oxygen reduction (Fig. 2A–D). Kadenbach’s theory postulates that stress removes ATP-dependent inhibition of CytOx [54], resulting in an increase in the ΔΨm and excessive formation of ROS [37]. However, the relationship between the ATP-dependent inhibition of CytOx enzyme activity, an increase of the ΔΨm and formation of ROS as a mitochondrial regulator is not yet known. We have observed that the rate of electron transmission on CytOx determines the inhibitory effect and the assumed subsequent production of ROS, which are generally considered a major cause of tissue damage [26, 27]. Thus, degenerative diseases and ageing could be better understood as an elementary mitochondrial process. Although debatable, phosphodiesterase inhibitors appear as key regulatory factors that influence the respiratory activity of CytOx. It is likely that analysis of PDE action can provide a framework for further studies because a variety of chemical compounds may affect oxidative phosphorylation to a much larger extent [122].

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