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Luca G Iacovino, University of Pavia
Nicola Manzella, Université de Toulouse
Jessica Resta, Université de Toulouse
Maria Antonietta Vanoni, University of Milan
Laura Rotilio, Univ Bari Aldo Moro
Leonardo Pisani, Emory University
Dale Edmondson, Emory University
Angelo Parini, Université de Toulouse
Andrea Mattevi, University of Pavia
Jeanne Mialet-Perez, Université de Toulouse

Only first 10 authors above; see publication for full author list.

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Rational Redesign of Monoamine Oxidase A into a Dehydrogenase to Probe ROS in Cardiac Aging

Luca Giacinto Iacovino, Nicola Manzella, Jessica Resta, Maria Antonietta Vanoni, Laura Rotilio, Leonardo Pisani, Dale Edward Edmondson, Angelo Parini, Andrea Mattevi, Jeanne Mialet-Perez, and Claudia Binda*

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ABSTRACT: Cardiac senescence is a typical chronic frailty condition in the elderly population, and cellular aging is often associated with oxidative stress. The mitochondrial-membrane flavoenzyme monoamine oxidase A (MAO A) catalyzes the oxidative deamination of neurotransmitters, and its expression increases in aged hearts. We produced recombinant human MAO A variants at Lys305 that play a key role in O2 reactivity leading to H2O2 production. The K305Q variant is as active as the wild-type enzyme, whereas K305M and K305S have 200-fold and 100-fold lower $k_{cat}$ values and similar $K_m$. Under anaerobic conditions, K305M MAO A was normally reduced by substrate, whereas reoxidation by O2 was much slower but could be accomplished by quinone electron acceptors. When overexpressed in cardiomyoblasts by adenoviral vectors, the K305M variant showed enzymatic turnover similar to that of the wild-type but displayed decreased ROS levels and senescence markers. These results might translate into pharmacological treatments as MAO inhibitors may attenuate cardiomyocytes aging.

Oxidative stress is a pathophysiological condition resulting from an imbalance of cellular free radicals (reactive oxygen or nitrogen species, ROS or RNS, respectively) with respect to antioxidant defense systems. This process is associated with a number of diseases such as cancer and neurodegenerative disorders, though the underlying molecular mechanisms are not fully understood. Many pathological or frailty states may lead to oxidative stress, but in certain circumstances ROS generated by cell oxidative metabolism may contribute themselves to triggering a disease. In this context, the role of H$_2$O$_2$ is dual because, while being physiologically involved in many signaling pathways, it represents a deleterious source of ROS through the Fenton or Haber−Weiss reaction when produced in excess.

Mitochondrial respiration is the main source of superoxide anions and H$_2$O$_2$, but the membrane of these organelles contains also other enzymes that generate ROS. Monoamine oxidases A and B (MAO A and MAO B, respectively) are 60-kDa mammalian flavoproteins that feature a water-soluble globular main body anchored to the outer mitochondrial membrane through a C-terminal α-helix. They belong to the FAD-dependent amine oxidase family of enzymes that catalyzes the oxidation of a carbon−nitrogen bond in various substrates with the concomitant reduction of the flavin coenzyme, which can be readily reoxidized by molecular oxygen leading to the generation of H$_2$O$_2$ (Figure 1). MAOs are particularly abundant in mammalian cells where they play a key role in the metabolism of both endogenous and exogenous neuroactive aromatic amines. The function of MAOs has been thoroughly studied in the central nervous system where they represent established drug targets for Parkinson’s disease and depression. However, MAOs are widely expressed also in non-neuronal tissues, including the heart, in which they regulate the local concentrations of serotonin, noradrenaline, and dopamine. Interestingly, MAO A levels in the heart increase significantly with aging. As a mimicry model of the elderly heart, the effects of cardiac overexpression and activation of MAO A in transgenic mice were studied by the authors of the present work, showing that an excess of H$_2$O$_2$ has multiple p53-mediated cell responses including mitochondrial damage and cell death. More recently, they also demonstrated that a chronic MAO A-dependent increase of H$_2$O$_2$ in heart cell lines triggers lipid peroxidation, elevated p53/p21 levels, DNA

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damage response, and other classical senescence markers. This cascade of events is associated with mitochondrial dysfunction, decreased respiration, and inhibition of parkin-mediated mitophagy. These findings have important clinical implications because MAO A-dependent release of H2O2 may occur either under acute pathological conditions such as ischemia-reperfusion injury when MAO substrates are heavily released or chronically when MAO A expression increases during aging.

Although we supported evidence for the involvement of H2O2 in acute and chronic cardiac damage, it is still uncertain whether MAO A-dependent substrate regulation may also have an impact on cardiac dysfunction. In an effort to probe the molecular features of human MAO A in relation to H2O2 production, we produced recombinant enzyme variants at the Lys305 site lying close to the flavin cofactor redox states are highlighted in black and yellow, respectively. In oxidases, the presence of a positive charge in proximity of the flavin isoalloxazine ring is believed to promote O2 binding and activation followed by H2O2 generation. In flavin-dependent amine oxidases, the conserved Lys305 lying on the top rim of the enzyme active site is H-bonded (dashed lines) to the flavin N5 atom through a bridging water molecule, which is believed to represent the O2 binding site.

The two reductive and oxidative half reactions related to the flavin cofactor redox states are highlighted in black and yellow, respectively. In oxidases, the presence of a positive charge in proximity of the flavin isoalloxazine ring is believed to promote O2 binding and activation followed by H2O2 generation. In flavin-dependent amine oxidases, the conserved Lys305 lying on the top rim of the enzyme active site is H-bonded (dashed lines) to the flavin N5 atom through a bridging water molecule, which is believed to represent the O2 binding site.

Figure 1. Scheme of amine substrate oxidation catalyzed by human MAO A (gray oval; the FAD coenzyme is covalently bound to a Cys residue). Oxidation of serotonin to 5-hydroxyindoleacetaldehyde (5-HIAL) is shown, with serotonin being the main MAO A substrate in heart metabolism. The two reductive and oxidative half reactions related to the flavin cofactor redox states are highlighted in black and yellow, respectively. In oxidases, the presence of a positive charge in proximity of the flavin isoalloxazine ring is believed to promote O2 binding and activation followed by H2O2 generation. In flavin-dependent amine oxidases, the conserved Lys305 lying on the top rim of the enzyme active site is H-bonded (dashed lines) to the flavin N5 atom through a bridging water molecule, which is believed to represent the O2 binding site.

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MAO covalent inhibitors are known that, though differing for the inactivation mechanism, all rely on the enzymatic functionality of the protein and the associated flavin cofactor. We exploited this property of MAOs to further test the Lys305 mutants. Within the propargylic class of molecules, clorgyline is a very well-known MAO A selective inhibitor that promptly forms an adduct with the flavin N5 atom that produces a sharp and intense peak at 415 nm in the UV−vis spectrum (Supporting Information Figure 1). Tranylcypromine and hydrazines are nonselective inhibitors of MAOs that behave as suicide substrates (i.e., implying a C−N bond oxidation) by reacting with either C4A or N5 atoms of the FAD cofactor, which can be detected spectrophotometrically as a bleached flavin peak. Though to different extents, both tranylcypromine and different hydrazine analogs are known to involve enzyme turnover (with molecular oxygen consumption) as part of the inactivation process; i.e., formation of the covalent adduct is preceded by unproductive cycles with a release of the oxidized inhibitor without formation of any bond with the flavin cofactor. Instead, inhibition by propargylamine compounds is not strictly dependent on the redox catalytic cycle of MAOs, being nevertheless affected by a lower functionality of the protein (as for K305M and K305S mutants).

| MAO A  | $k_{cat}$ (min$^{-1}$) | $K_m$ (mM) | $k_{cat}/K_m$ (min$^{-1}$mM$^{-1}$) |
|--------|----------------------|-----------|-----------------------------------|
| MMTP direct assay ($\varepsilon_{420} = 25000$ M$^{-1}$ cm$^{-1}$) |
| wild-type | 51.60 ± 1.70 | 0.18 ± 0.02 | 286.66 |
| K305M | 0.28 ± 0.01 | 0.36 ± 0.05 | 0.77 |
| K305S | 0.53 ± 0.03 | 0.20 ± 0.04 | 2.65 |
| K305Q | 21.4 ± 0.54 | 0.04 ± 0.005 | 528.50 |
| K305R | not active | not active | not active |
| kynuramine direct assay ($\varepsilon_{316} = 12000$ M$^{-1}$ cm$^{-1}$) |
| wild-type | 120.20 ± 6.70 | 0.15 ± 0.01 | 801 |
| K305M | 0.67 ± 0.02 | 0.18 ± 0.02 | 3.72 |
| K305S | 0.98 ± 0.03 | 0.18 ± 0.03 | 5.44 |
| K305Q | 91.39 ± 2.81 | 0.16 ± 0.02 | 571.18 |
| K305R | not active | not active | not active |

“All details related to the experiments are reported in the Supporting Information. Briefly, all assays were performed at 25 °C in 50 mM HEPES/NaOH at pH 7.5 containing 0.25% reduced Triton X-100 (air saturated solution). Enzyme concentration was 0.07 μM for MAO A wt and 1.8 μM for MAO A mutants.

Figure 2. Spectrophotometric measurements of MAO A activity under anaerobiosis conditions. In all experiments, the cuvette contained 10 μM enzyme in 50 mM potassium phosphate at pH 7.8, 300 mM sodium chloride, 20% (w/v) glycerol, and 0.05% (w/v) Fox-Choline-12. (A) Flavin reduction was obtained by anaerobically adding 1 mM tyramine; reoxidation by molecular oxygen was monitored for the K305M mutant (red) compared to the wild-type enzyme (black). Enzyme reduction (left panel) was followed by measuring the absorbance at 456 nm corresponding to the peak of the oxidized flavin spectrum, which is bleached when flavin is reduced by the amine substrate. Enzyme reoxidation was monitored through the reappearance of the peak centered at 450 nm after exposure of the reaction mix to oxygen. Supporting Information Figure 2A and B show the overall UV−vis spectra of the oxidized and reduced enzyme for wild-type and K305M, respectively. (B) K305M flavin reoxidation by alternative electron acceptors: 200 μM benzoquinone (left) and 50 μM coenzyme Q$_0$ (right). UV−vis spectra of the oxidized (initial), photoreduced, and reoxidized K305M mutant are depicted as continuous black, dashed black, and gray lines, respectively. In this experiment, photochemical reduction of the enzyme was preferred to avoid multiple turnovers. In both panels, the profile of the photoreduced enzyme is consistent with a mixture of the anionic semiquinone and hydroquinone flavin forms that were previously observed for MAOs. The peak at 350 nm in the left panel is due to benzoquinone absorbance.
measurements showed that the K305Q can be inactivated to the same extent as the wild-type enzyme by all inhibitors, whereas inactivation of both K305M and K305S is far less effective (Supporting Information Figure 1). Furthermore, these two mutants react less efficiently with those inhibitors that require enzyme turnover such as tranylcypromine and phenelzine. The only exception is represented by phenylhydrazine that takes about 20 min to inactivate all proteins, most likely due to the lower oxygen consumption required for the inactivation process.

The experiments described above showed that the K305M mutant is impaired in enzymatic activity, but it is not completely inactive (as for K305R). MAO A catalyzes a redox reaction in which the amine substrate is oxidized with the concomitant reduction of the FAD cofactor (reductive half reaction; Figure 1). The catalytic turnover is guaranteed by the prompt flavin reoxidation by molecular O$_2$ (oxidative half reaction).4,17 To better dissect the role of Lys305 in H$_2$O$_2$ production by MAO A, we monitored the K305M enzyme reaction spectrophotometrically under anaerobiosis conditions, which were essential to keep the enzyme in the reduced state as the mutant is likely to retain some O$_2$ reactivity. This specific mutant (rather than K305S) was selected for these further studies because its side chain structurally mimics that of lysine, though returning an impaired turnover efficiency due to the lack of the positive charge that prevents an optimal interaction with the flavin through water as observed in wild-type MAO A. Incubation of the oxygen-free enzyme sample with a tyramine substrate led to full reduction of the flavin coenzyme, which could be stably reached with both wild-type and the K305M mutant in a few minutes (Figure 2A). When aerobic conditions were restored by exposing the system to air, the wild-type enzyme was promptly reoxidized, whereas in the case of K305M the reappearance of the oxidized flavin peak was only gradually obtained and required more than 1 h to reach completion. In agreement with the steady-state kinetic parameters (Table 1), these results indicate that K305M can bind and oxidize the amine substrate like the wild-type enzyme, while it is heavily impaired in oxidized flavin regeneration by molecular oxygen.

Next, we asked if the K305M mutant could be reoxidized by electron acceptors alternative to molecular oxygen. Although MAOs are “true” oxidases, their active site might bind organic molecules that accept electrons and, in the case of the K305M mutant, may work more efficiently than oxygen. Among these, quinones are electrophilic compounds normally found in mitochondria and other organelles where they serve as acceptors in cellular electron transport chains. It was reported that some quinones and other redox compounds display binding affinity for MAO A.19,20 We tested benzoquinone and coenzyme Q$_0$ for their ability to reoxidize the K305M mutant under anaerobiosis conditions (Figure 2B).
basic unit of quinones, the latter is the reactive part of the naturally occurring coenzyme Q₉₉ that is highly hydrophobic and therefore very difficult to experimentally handle in aqueous solutions. For this experiment, K305M MAO A was anaerobically photoreduced rather than incubated with substrate to avoid any direct electron transfer from the substrate to the electron acceptor or multiple turnovers. The addition of either benzoquinone or coenzyme Q₀ readily led to the full reappearance of the peak at 456 nm, indicating that these molecules can reoxidize the enzyme-bound FAD coenzyme. The same experiment was carried out also with the wild-type enzyme using benzoquinone (Supporting Information Figure 3) obtaining similar results. Benzoquinone and coenzyme Q₀ were also tested for their ability to restore enzymatic activity of K305M was observed. This may be due to the fact that the mutant is endowed with, though limited, reactivity with oxygen and that the large amount of substrate may hamper the accessibility of the alternative electron acceptor. These results outlined the K305M MAO A mutant as an oxygen-inert tool to be tested in a cellular model of cardiomyocytes where the bioavailability of quinone-based electron acceptors may restore an efficient enzyme regeneration.

We thereby generated two adenoviral constructs in order to overexpress wild-type (Ad WT) and K305M (Ad K305M) human MAO A in cardiomyoblasts. As shown in Figure 3A, MAO A protein levels were similarly increased in cells transduced with Ad WT or Ad K305M, compared to control untransduced cells (Ctr). Interestingly, MAO A activity, measured as the quantity of products formed by ¹⁴C-S-HT degradation, was equally increased in Ad WT and Ad K305M-transduced cells compared to Ctr. Thus, K305M appears to have turnover activity similarly to wild-type when expressed in cells, which raised the question whether the mutant uses O₂ as an alternative electron acceptor in cells. We directly measured the concentration of H₂O₂ generated during the oxidation of tyramine. As shown in Figure 3B, tyramine addition led to H₂O₂ production in Ctr cells with a maximum at 60 min, corresponding to endogenous MAO A activation. Most interestingly, Ad WT transduction potentiated H₂O₂ production upon tyramine administration, which was instead not observed for Ad K305M. This demonstrates that K305M mutant is able to degrade MAO substrate without production of H₂O₂. Thus, we took advantage of this unique property to establish the contribution of ROS in the senescent response induced by MAO A. To better establish the specific effects of wild-type and K305M MAO A in cardiomyoblasts, we blocked endogenous MAO A with the irreversible inhibitor clorgyline before performing adenoviral transduction and treatment with tyramine (Supporting Information Figure 4A). In cells transduced with Ad WT, but not K305M, tyramine application for 30 min significantly increased ROS production compared to Ctr (Supporting Information Figure 4). We next evaluated the chronic effect of MAO A stimulation over 3 days with tyramine on aging markers by measuring senescence-associated-β-gal (SA-β-gal), a β-galactosidase activity detectable at pH 6.0 in senescent cells. Interestingly, SA-β-gal activity was increased only in cells expressing Ad WT but not K305M. Similarly, the senescence marker p21 accumulated in Ad WT cells but not in Ad K305M cells, while the retinoblastoma protein (Rb) was dephosphorylated only in tyramine-activated Ad WT cells, preventing the progression of the cell cycle (Figure 3D). Altogether, our results show that substituting Lys305 with Met in MAO A impairs the production of H₂O₂, preventing oxidative stress and senescence induced by a chronic activation of MAO A.

In conclusion, our work provided an insightful investigation on the role of the conserved lysine residue of flavin-dependent amine oxidases in enzyme turnover and on the effects of H₂O₂ generated by human MAO A in cardiac cell aging. We demonstrated that in human MAO A, the K305M mutant retains the capability to bind and oxidize the amine substrate, while it is significantly impaired in reoxidation by molecular O₂. Using this mutant as a mimic of an oxygen-inert enzyme in the context of cardiomyocyte cells, we gave further support to the hypothesis that, in the heart, MAO A represents a noteworthy source of ROS promoting cell senescence. From this perspective, pharmacological treatments targeting MAO A, so far limited to neurological diseases, may be extended to prevent heart failure in some frailty conditions of aged patients. Moreover, since many reports indicate that MAO A promotes prostate and glioblastoma tumorigenesis and metastasis, this approach may also limit cancer growth.21

**ASSOCIATED CONTENT**

**Supporting Information**

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acschembio.0c00366.

Details of the experimental methods, supporting Table 1, supporting Figures 1−4 (PDF)

**AUTHOR INFORMATION**

**Corresponding Author**

Claudia Binda − Department of Biology and Biotechnology, University of Pavia, Milan, Italy; orcid.org/0000-0003-2038-9845; Phone: (+39)0382-985527; Email: claudia.binda@unipv.it

**Authors**

Luca Giacinto Iacovino − Department of Biology and Biotechnology, University of Pavia, Milan, Italy
Nicola Manzella − Institute of Metabolic and Cardiovascular Diseases (I2MC), Institut National de la Sante et de la Recherche Medicale (INSERM), Université de Toulouse, Toulouse, France
Jessica Resta − Institute of Metabolic and Cardiovascular Diseases (I2MC), Institut National de la Sante et de la Recherche Medicale (INSERM), Université de Toulouse, Toulouse, France
Maria Antonietta Vanoni − Department of Biosciences, University of Milan, Milan, Italy; orcid.org/0000-0001-7213-732X
Laura Rotilio − Department of Biology and Biotechnology, University of Pavia, Milan, Italy
Leonardo Pisani − Department of Pharmacy-Drug Sciences, University of Bari Aldo Moro, Bari, Italy; orcid.org/0000-0002-4198-3897
Dale Edward Edmondson − Department of Biochemistry, Emory University, Atlanta, Georgia, United States; orcid.org/0000-0002-1304-231X
Angelo Parini − Institute of Metabolic and Cardiovascular Diseases (I2MC), Institut National de la Sante et de la
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Notes

The authors declare no competing financial interest.

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