Sabotage at the Powerhouse? Unraveling the Molecular Target of 2-Isopropylbenzaldehyde Thiosemicarbazone, a Specific Inhibitor of Aflatoxin Biosynthesis and Sclerotia Development in Aspergillus flavus, Using Yeast as a Model System

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Abstract: Amongst the various approaches to contain aflatoxin contamination of feed and food commodities, the use of inhibitors of fungal growth and/or toxin biosynthesis is showing great promise for the implementation or the replacement of conventional pesticide-based strategies. Several inhibition mechanisms were found taking place at different levels in the biology of the aflatoxin-producing fungal species such as Aspergillus flavus: compounds that influence aflatoxin production may block the biosynthetic pathway through the direct control of genes belonging to the aflatoxin gene cluster, or interfere with one or more of the several steps involved in the aflatoxin metabolism upstream. Recent findings pointed to mitochondrial functionality as one of the potential targets of some aflatoxin inhibitors. Additionally, we have recently reported that the effect of a compound belonging to the class of thiosemicarbazones might be related to the energy generation/carbon flow and redox homeostasis control by the fungal cell. Here, we report our investigation about a putative molecular target of the 3-isopropylbenzaldehyde thiosemicarbazone (mHtcum), using the yeast Saccharomyces cerevisiae as model system, to demonstrate how the compound can actually interfere with the mitochondrial respiratory chain.

Keywords: aflatoxin inhibitor; mitochondrion; respiratory chain; yeast model system

1. Introduction

Aflatoxin (AF) contamination of food and feed commodities is a major health concern for sanitary authorities all over the world, since these mycotoxins represent a health hazard to both animals and humans [1]. In particular, aflatoxin B1 is considered to be one of the most carcinogenic natural products. Aspergillus flavus and Aspergillus parasiticus are the main producers of this type of mycotoxin, and both fungi have a wide geographic distribution and may contaminate agricultural crops important for animal and human diet [2]. Moreover, global climate changes have exacerbated the diffusion of these mycotoxigenic Aspergilli in regions that, until recently, were considered to be temperate climate areas and were not prone to their growth and their overwinter persistence [3,4].

AF contamination may occur pre-harvest or post-harvest owing to fungal infection of crops in the fields or during storage. Accordingly, different approaches have been developed to counteract mycelia and/or mycotoxin contamination. These include good agronomic practices to prevent plant stress that
may weaken plant defense or stimulate mycotoxins biosynthesis by the fungus, or chemical treatments
to avoid damage of kernels by insects and bio-control by using natural competitors to displace the
threatening organism from the ecological niche [3]. Additionally, AF is highly persistent on the various
food matrices and is scarcely degraded by the industrial transformation procedures. Carryover of AFs
along the food chain may be the primary cause of acute toxicosis in humans and animals, however
the possibility that the Aspergillus infection of tissues, organs etc., which is generally referred to
as aspergillosis, may lead to AF production “in situ” (i.e. in the affected body district) should not
be discarded [5–9]. Moreover, mycotoxin production may increase the virulence of aspergillosis by
depressing the immunological response of the organism [10]. The availability of specific inhibitors of
AF biosynthesis may thus be useful not only in an agronomic perspective, but also as a weapon to
counteract the consequences of fungal infection in humans and animals.

An impressive amount of data is reported in the literature concerning the fungistatic and anti-toxin
activity of natural derived and synthetic compounds. However, the molecular mechanisms involved
in the specific inhibition of AF synthesis by the relevant compound remain rather speculative in many
cases, even if several metabolic/regulatory targets have been suggested [11–31]. At present, we are
facing a general consensus that AF has a prominent role in the oxidative stress response by the fungus
even if, due to the complexity of the regulatory and metabolic steps that are involved in the response,
several details of the framework are still unavailable.

We have recently described a set of molecules belonging to the class of thiosemicarbazones,
that were tested for their fungicidal and/or inhibitory activity on mycotoxin accumulation on cereal
crops [32–34]. These molecules, which are very attractive for their coordinating versatility and have the
possibility to easily modify the molecular backbone and tune their physical and chemical properties,
were designed in order to take advantage of the individual antymycotic properties of metal ions (such
as copper, for instance) and natural aldehydes contained in spices are known for their antymycotic
properties. For this scope, the aldehydes were functionalized with thiosemicarbazone to endow
them with chelating properties towards metal ions. Unexpectedly, these intermediates also showed
interesting properties. In particular one of the most promising of them, the thiosemicarbazone of
the meta-isomer of cuminaldehyde, namely isopropylbenzaldehyde thiosemicarbazone (mHtcum),
displayed a specific activity against aflatoxin biosynthesis distinct from a fungistatic effect, and was
therefore investigated in depth to help understand its mode of action in the fungal cell [33].

The data gathered by the above-mentioned experiments were consistent with a possible molecular
target involved in the energy generation/carbon flow and the redox homeostasis control by the cell: in
particular, some evidence suggested that mHtcum could interfere at the mitochondrial level. Here,
we describe a set of experiments performed in the yeast Saccharomyces cerevisiae addressed to the
characterization of mHtcum effect and to the identification of its possible molecular target(s). Even
though yeast does not possess the secondary metabolism pathways involved in aflatoxin synthesis,
it shares all basal pathways for energy production with other fungi and will be used as a model to
corroborate our hypothesis.

2. Results and Discussion
2.1. Effect of mHtcum on the Oxidative Carbon Source Utilization

On the basis of genetics as well as biochemical and proteomic data, it has been previously
speculated that isopropylbenzaldehyde thiosemicarbazone (mHtcum) may negatively affect aflatoxin
biosynthesis in Aspergillus flavus by redirecting carbon flow in the cell and by modulating the activity
of enzymes involved in energy metabolism [33]. In order to investigate whether mHtcum activity may
be coupled to a switch from fermentative to respiratory metabolism (or vice versa), we first performed a
series of experiment using the yeast S. cerevisiae as a model system. We analyzed the effect of mHtcum
on the utilization of fermentable (glucose) and non-fermentable (ethanol) carbon source by performing
a disk diffusion assay, as reported in Figure 1. The effect of other thiosemicarbazones, previously described for their antifungal and antiaflatoxigenic effect [32], was also compared.

On glucose, no growth inhibition was observed for all the compounds. In contrast, in an ethanol-containing medium, an inhibition halo was observed in the case of Htcin, Htcum, mHtcum and oHtcum, i.e. the molecules that inhibit aflatoxin biosynthesis in A. flavus at the highest level. Similar results were obtained when ethanol was replaced with other non-fermentable carbon sources such as glycerol and acetate (Figure S1, Supplementary Material), leading us to exclude the possibility of an ethanol-specific effect.

In addition, the response of S. cerevisiae to the mHtcum antiaflatoxigenic concentration of 50 μM during a non-fermentable metabolism was tested with a spot assay (Figure 2); an inhibitory effect of the thiosemicarbazone on yeast cell proliferation was detectable at the 10⁴ cells/spot concentration, and dramatically increased as the cell concentration decreased.

2.2. Interference of mHtcum on Mitochondrial Activity

The observation that mHtcum negatively affects yeast growth only in the presence of an oxidative carbon source points to a possible molecule-induced mitochondrial impairment. Thus, we wondered if mHtcum might interfere with mitochondrial respiratory-linked processes in S. cerevisiae, in particular
with the respiratory activity and the mitochondrial cytochromes content, but also with gametogenesis. In fact, nutrient sensing, and in particular the carbon/energy status, plays a critical role in yeast sporulation, and various respiratory inhibitors are reported to affect this developmental process [35,36].

For the yeast sporification assay, the diploid strain W303, pre-grown for five days in YP medium, was incubated in the sporification medium added with mHtcum 25, 50 and 100 μM. Asci production was estimated after 7 days; antimycin (AMY), a specific inhibitor of the mitochondrial respiratory chain, was used as a reference. As expected, AMY highly affected the number of asci; at the same time, mHtcum was highly effective in lowering yeast sporulation in a dose-dependent manner (Figure 3A).

![Image](image-url)

**Figure 3.** Interference of mHtcum on mitochondrial activity in *S. cerevisiae*. (A) Asci production in W303 diploid strain treated with mHtcum. The effect of thiosemicarbazone was compared with the effect of the mitochondrial respiratory inhibitor antimycin A (AMY). Values are expressed as percentage of ascis with respect to the number of total cells; statistically significant differences from the control (CNT) were indicated with an asterisk (*p* ≤ 0.05). (B) Oxygen consumption rate. W303-1B grown in the absence (CNT) or in the presence of mHtcum at different concentrations (from 5 to 50 μM). Values were normalized to the untreated strain and represented as the mean of at least three values ± SD. Values significantly different from CNT were indicated with an asterisk (*p* ≤ 0.05). (C) Reduced versus oxidized cytochrome spectra: peaks at 550, 560 and 602 nm correspond to cytochromes c, b and aa3, respectively. The height of each peak relative to the baseline is an index of cytochrome content. (D) Mitochondrial DNA mutability. Frequency of respiratory deficient (*petite*) cells determined in untreated (CNT) or 50 μM mHtcum treated yeast cultures, expressed as a percentage.

The effect of mHtcum treatment on oxygen consumption was then measured in yeast cells pre-grown for 18 hours in the presence of different concentrations of mHtcum. As reported in Figure 3B, mHtcum thiosemicarbazone caused a concentration-dependent decrease of respiration, reaching a 55% of inhibition at 50 μM. Accordingly with the drastic impairment of the respiration rate, the profile of mitochondrial cytochromes of yeast cultures was negatively affected by mHtcum treatment in a concentration-dependent manner (Figure 3C).

The decrease of cell respiration and of the number of cytochromes might be due to the inhibition of at least one of the steps of the electron flow along the respiratory chain by mHtcum. On the other hand, since proteins belonging to the respiratory chain are encoded both by nuclear and mitochondrial genes, the alteration of OXPHOS metabolism in the presence of mHtcum might be due to a mutagenic effect on mitochondrial DNA, with a consequent absence of mitochondrial encoded respiratory proteins. In order to discriminate between these two hypothesis, we measured the frequency of cytoplasmic
mitochondrial mutants (petite mutants) displaying a respiratory deficient phenotype after treatment with mHtcum. Data reported in Figure 3D show that treatment with thiosemicarbazone of the wild type yeast strain W303-1B did not increase mitochondrial DNA mutability in terms of the petite mutants ratio. Results are consistent with our previous data suggesting that mHtcum exerts a sort of rescue effect on petite high-frequency yeast mutants [33], owing at least in part to its antioxidant activity.

To evaluate if the long-term effect of mHtcum in yeast cells during growth may be exerted at the level of respiratory complex proteins, the relative abundance of COR1 and COR2 (core subunit 1 and subunit 2 respectively of the ubiquinol-cytochrome c reductase; complex III), COX2 (subunit II of cytochrome c oxidase; complex IV) and ATP2 (beta subunit of the F1 sector of mitochondrial F1F0 ATP synthase; complex V) was estimated in the same treated cultures used for the oxygen consumption rate and discussed above.

Densitometric analysis of immunoreactive signals revealed that COR1 and COR2 level was significantly decreased by mHtcum treatment with respect to the control, while the level of COX2 was not significantly affected (Figure 4). Even more remarkably, an increase of ATP2 in yeast cells was detected: the up-regulation of the beta subunit of mitochondrial ATP synthase was in fact observed also in A. flavus cultures treated with mHtcum, as described in Degola et al. [33].

The effect of the thiosemicarbazone on the enzymatic activity of complexes II, III and IV was then evaluated: an enriched fraction of mitochondria extracted from yeast cultures was used for the spectrophotometric assays, in which the thiosemicarbazone was added at increasing concentrations (from 50 to 150 μM) and compared with the relevant specific inhibitors (malonate, AMY and Na-azide respectively). A negative effect of mHtcum was observed on the complex III, whose specific activity decreased by 25–30% at the higher thiosemicarbazone concentrations (Figure 5). In contrast, complex II and IV activities seemed to be insensitive to the addition of mHtcum at any concentration tested (Figure 5). This result supports the hypothesis of a defect in the electron flow along the respiratory chain induced by mHtcum, suggesting, in particular, a specific inhibition of complex III.
We wondered, therefore, if mHtcum inhibition of complex III might be due to its ability to compete with the aromatic ring of mHtcum can be easily stabilized. In addition, a strong hydrogen bond (2.001 Å) between thiosemicarbazone N2 and the carboxyl group of Asp229 is predicted: this bond may convincingly be responsible for the maintenance of mHtcum into the binding site, facing the aromatic portion of the ligand with the hydrophobic cavity of the protein. Noticeably, as for AMY, mHtcum resulted close to heme b1 (3.376 Å); this proximity is considered to be fundamental for the inhibition mechanism of AMY, since the b1 heme is involved in the oxidation of ubiquinol to ubiquinone, a crucial step in the Q-cycle.

2.3. Comparative Molecular Docking Analysis

The most studied inhibitor of complex III is Antimycin A, that interacts with the Qi/Qn site of cytochrome c reductase in mitochondrial complex III [37], causing a collapse of the proton gradient across the mitochondrial inner membrane, leading to the loss of the membrane potential and, as a consequence, an increase of ROS production and a reduction of the ATP intracellular level [38,39]. We wondered, therefore, if mHtcum inhibition of complex III might be due to its ability to compete with the substrate at Qi site, interfering with the electron flow. Thus, a structure-based docking study was performed to investigate the intermolecular interactions between our thiosemicarbazone and the mitochondrial complex III. Since the inhibitory mechanism of AMY is caused by its binding at bc1cytochrome level in the Qi site, resulting in the disruption of the Q-cycle and the entire enzyme turnover, the docking method was developed using GOLD v5.5 and optimized to compare the theoretical positioning of mHtcum compared to AMY. Structural information about the protein was taken from the Protein Data Bank accession 3BCC (10.2210/pdb3BCC/pdb), antimycin bound to the cytochrome bc1 complex from chicken with AMY, as co-crystallized inhibitor, located in the Qi region. The protein model was prepared by deleting solvent molecules and AMY from the structure and by adding the hydrogen atoms. A 6Å-cavity centered in the Qi region of the protein was used as binding site. A 50-run docking approach was used and the results obtained were scored using the Chemscore fitness function. All the poses were then rescored using the ChemPLP function. The entire docking protocol was validated using the self-docking approach, and AMY was docked successfully [40,41], with RMSD 1.403 Å.

Figure 6 shows the best pose of mHtcum into the Qi site: in detail, the hydrophobic environment formed by Phe221, Ile228, Tyr225, Ala24, Trp32 and Leu201 [42] allows the formation of a binding pocket in which the aromatic ring of mHtcum can be efficiently stabilized. In addition, a strong hydrogen bond (2.001 Å) between thiosemicarbazone N2 and the carboxyl group of Asp229 is predicted: this bond may convincingly be responsible for the maintenance of mHtcum into the binding site, facing the aromatic portion of the ligand with the hydrophobic cavity of the protein. Noticeably, as for AMY, mHtcum resulted close to heme b1 (3.376 Å); this proximity is considered to be fundamental for the inhibition mechanism of AMY, since the b1 heme is involved in the oxidation of ubiquinol to ubiquinone, a crucial step in the Q-cycle.
was used as a reference to evaluate the binding strength of mHtcum in the Qi region. Very similar binding energies were obtained for thiosemicarbazone and AMY, $\Delta G_{\text{bind}} = -5.96$ kcal/mol and $\Delta G_{\text{bind}} = -6.05$ kcal/mol respectively, suggesting that mHtcum is able to bind at Qi site thus decreasing the activity of mitochondrial complex III.

2.4. A. Flavus Growth, Development and Gene Expression

Recently, Sakuda and coworkers [23] have shown that different respiration inhibitors such as Na-azide, rotenone, oligomycin and antimycin A, that specifically affect different steps of the respiratory chain, inhibit the biosynthesis of aflatoxin in A. flavus without significantly interfere with fungal growth. However, no correlation between the inhibitor target along the respiratory chain and the level of inhibition activity on the toxin accumulation have been demonstrated, suggesting that a block of respiration “tout court” may be the cause of the inhibition of aflatoxin biosynthesis. Due to the similar behavior observed for mHtcum, we wondered if the compound may mimic the activity of one of the natural inhibitors of respiration reported above.

All previously discussed evidence seemed to confirm the hypothesis that in yeast cells a common molecular target, or probably one of many, might be shared by mHtcum and AMY, and that this target is located at the mitochondrial level, controlling the functionality of the respiratory chain. We then compared the effect of 50 $\mu$M antimycin and mHtcum on A. flavus; the fungus radial growth was evaluated on a fermentable (2% glucose containing medium) and a non-fermentable carbon source (2% ethanol containing medium). Similarly to what was observed in sucrose fed cultures, in which the thiosemicarbazone did not significantly affect either conidia germination or mycelium growth [33], the A. flavus colonies inhibition rate in glucose was lower than 5% with respect to the control. In contrast, in the presence of a non-fermentable carbon source, mHtcum exposure led to a 18% of radial growth inhibition. Even at a higher level, AMY treated cultures showed the same behavior (Figure 7A).
The response of *A. flavus* in terms of aflatoxin accumulation and sclerotia biogenesis when treated with AMY or mHtcum was additionally evaluated in order to compare the effect of these molecules on physiological processes belonging to the fungus secondary metabolism. Consistently with Sakuda et al. [23] antimycin A was effective in lowering aflatoxin accumulation at an inhibition level comparable to mHtcum (up to 75% and 70% respectively). In contrast, the sclerotia production was not affected by the presence of AMY, while the *A. flavus* cultures showed to be highly sensitive to the treatment with the thiosemicarbazone (up to 85% of sclerotia inhibition with respect to the control; Figure 7B).

An additional element of the differential activity exerted by the two compounds was then obtained through the analysis of specific gene expression: surprisingly *AflR*, the major regulator of the aflatoxin gene cluster, and the *aflD* and *omtB* genes, encoding for two structural enzymes directly involved in the toxin biosynthesis, proved to be non-down-regulated by antimycin A, contrary to what was observed with the mHtcum treatment (Figure 7C). Similar results were previously reported by Sakuda et al. [23] for inhibitors of mitochondria complex I and II, reinforcing the hypothesis that the mechanism involved in the block of aflatoxin production by antimycin A (and the above mentioned inhibitors) might be only in part shared (differently from the one triggered) by mHtcum. This, in fact, showed down-regulation of the expression of both *aflR* and the other two structural genes (*aflD* and *omtB*) (Figure 7C).
3. Materials and Methods

3.1. Thiosemicarbazones

Cuminaldehydethiosemicarbazone (Htcum), trans-cinnamaldehyde thiosemicarbazone (Htcin), quinoline-2-carboxyaldehyde thiosemicarbazone (Htisq), 5-fluoroisatin N4-methylthiosemicarbazone (HtmeFis) and 5-fluoroisatin thiosemicarbazone (HtFis) were previously described in Degola et al. [32]; 2-isopropylbenzaldehydethiosemicarbazone (mHtcum) and 3-isopropylbenzaldehyde thiosemicarbazone (oHtcum) were previously described in Degola et al. [33].

3.2. Microbial Strains, Media and Culture Conditions

Aspergillus flavus strains used in this work were isolated from corn fields of the Po Valley [45] and have already been used for previous experimentation concerning thiosemicarbazone activity [32,33]. A. flavus strains are available from the corresponding author on request. Conidia suspensions were obtained from 10-day YES-agar [2% \((w/v)\) yeast extract (Difco, Detroit, MI), 5% \((w/v)\) sucrose (Sigma, St Louis, MO), 2% \((w/v)\) agar (Difco)] cultures incubated at 28 °C; conidia concentration (quantified by \(OD_{600}\)) and viability (>90%) were determined according to Degola et al. [45]. Coconut milk-derived medium (CCM) used for microplate assays was obtained as described in Degola et al. [46].

Saccharomyces cerevisiae yeast strains used in this study were the haploid strain W303-1B (MAT\(\alpha\) ade2-1 leu2-3,112 ura3-1 trp1-1 his3-11,15 can1-100) [47] and the diploid strain W303 (a/\(\alpha\) ade2 leu2 ura3 trp1 his3). All experiments were performed in YP medium (1% yeast extract, 2% peptone, ForMedium). Media were supplemented with various carbon sources as indicated in the text, in liquid phase or after solidification with 20 g/L agar (ForMedium). Growth in the liquid phase was performed with constant shaking at 28 °C.

3.3. Agar Disk-Diffusion Testing

The response of yeast cells to different thiosemicarbazones onto fermentable and non-fermentable carbon sources was assessed in YP medium: W303-1B strain liquid culture was properly diluted to 2.5 \(\times\) 10\(^6\) cells/mL, then 400 \(\mu\)L were seeded on YP agar plates supplemented with 2% glucose, 2% ethanol, glycerol or 2% k-acetate alternatively. Filter paper discs (about 6 mm in diameter) were placed on the agar surface, then a volume of 4 \(\mu\)L of 10 mM stock solution of thiosemicarbazones dimethylsulphoxide-resuspended was dropped onto. An equal volume of dimethylsulphoxide (DMSO) was used as control. Petri dishes were incubated at 28 °C, and after 48 h the diameters of inhibition growth zones were evaluated.

3.4. Spot Assay for Yeast Oxidative Growth

W303-1B strain were serially diluted from \(10^7\) to \(10^2\) cells/mL, then 10 \(\mu\)L of each dilution was spotted on YP agar plates supplemented with 2% glucose or 2% ethanol, with or without 50 \(\mu\)M isopropylbenzaldehyde thiosemicarbazone (mHtcum), in order to obtain spots containing from \(10^5\) to \(10^0\) cell/spot. Plates were incubated at 28 °C.

3.5. Asci Production in S. cerevisiae

Diploid W303 yeast strain cells were pre-grown onto 2% glucose YP agar plates for three days at 28 °C. A small portion of culture was then picked up with a sterile loop and straight-inoculated on the surface of SPO IV medium (0.25% \((w/v)\) yeast extract, 2% \((w/v)\) K-acetate, 0.1% \((w/v)\) glucose) agar plates supplemented with thiosemicarbazone mHtcum at increasing concentration (from 25 to 100 \(\mu\)M). DMSO and antimycin A (AMY) were used as negative and positive controls, respectively. Cells were left for sporulation at 28 °C for 6 days, then a small amount of cells was sampled from each treatment, resuspended in bidistilled water, loaded on a slide and observed with an inverted microscope at 200× magnification: three samples from each treatment were observed, and asci from a
total of five fields for each samples were counted. Experiments were conducted in triplicate. Results were expressed as the percentage of asci respect with the number of total cells.

3.6. Mitochondrial Respiratory Activity and Cytochrome Profiles

Oxygen consumption rate was measured at 30 °C using a Clark-type oxygen electrode (Oxygraph System Hansatech Instruments), with 1 mL of air-saturated respiration buffer (0.1 M phthalate KOH, pH 5.0), 0.5% glucose. Yeast W303-1B cells were cultured at 28 °C in YP medium supplemented with glucose at the non-repressing concentration of 0.6%, until glucose exhaustion, in the presence of isopropylbenzaldehyde thiosemicarbazone (mHtcum) at different concentrations (5, 10, 25, and 50 µM). The control was treated with the same volume of the solvent DMSO. Oxygen consumption was normalized to the dry weight of the cells. Cytochrome profiles were determined spectrophotometrically at room temperature (Varian Cary300 UV-VIS Spectrophotometer), by recording reduced oxidized cytochrome spectra of yeast W303-1B cells cultured for oxygen consumption rate determination.

3.7. Mitochondrial DNA Stability

For petite frequency determination the W303-1B strain was pre-grown at 28 °C in YP medium, supplemented with 2% ethanol in order to counterselect the petite cells that were already present in the population. Cells were then inoculated in YP medium supplemented with 2% glucose and isopropylbenzaldehyde thiosemicarbazone at the final concentration of 50 µM, whereas to the control treatment the same volume of DMSO was added; cultures were then incubated at 28 °C. After 15 generations, cells were plated at a dilution which provided approximately 200 cells/plate on SC (0.19% YNB without aminoacids and NH4SO4 powder ForMedium, 0.5% NH4SO4, supplemented with 1g/L dropout mix [48]) agar plates containing 2% ethanol and 0.25% glucose. Petite frequency was defined as the percentage of colonies showing the petite phenotype after a 6-day incubation at 28 °C.

3.8. Isolation of Mitochondria, Gel Electrophoresis and Western Blot Analysis

Mitochondrial proteins for Western Blot analyses were obtained by suspending the cells in extraction buffer containing 0.6 M sorbitol, 10 mM imidazole, 0.5 mM EDTA, 0.1% BSA and 1 mM PMSF. Cells were broken by vortexing on ice using glass beads and mitochondrial proteins were obtained by centrifugation and re-suspended in the extraction buffer. Quantification of protein concentration was performed by Bradford’s method [49], using Bio-Rad protein assay following the manufacturer’s instructions. For each sample, 15 µg of mitochondrial total proteins were loaded on 12% polyacrylamide gel, subjected to SDS-PAGE, and Western Blot was performed. Gels were electroblotted onto nitrocellulose filters and successively immunostained with specific antibodies against CORE1 and CORE2 (kindly provided by Prof. B.L. Trumpower), COX2 (Molecular Probes, Invitrogen), and ATP2 (kindly provided by Prof. J.P. di Rago). POR1 (Abcam Mitoscience, Cambridge, UK) was used as the loading control. After incubation with the appropriate secondary antibodies, ECL Western Blotting Substrate (ClarityTM, BioRad, Hercules, CA, USA) was used for final detection. Signals were quantified through QuantityOne Software (Bio-Rad, Hercules, CA, USA).

Enzymatic activities: Complex II (succinate dehydrogenase), complex III (cytochrome c reductase) and complex IV (cytochrome c oxidase) specific activities were measured spectrophotometrically according to Barrientos et al. [50] on a mitochondrial-enriched fraction prepared as previously described [51]. Thiosemicarbazone mHtcum was added to the cuvette. As control the same volume of DMSO was added.

3.9. Aflatoxin Production Assay

Effect on aflatoxin biosynthesis was assessed by the microplate fluorescence-based procedure described in Degola et al. [46]. Standard flat-bottom 96-well microplates (Sarstedt, Newton, NC, USA) were used. Suspensions of conidia were diluted and brought to the final concentration of 5 × 10^5 conidia/well; cultures were set in a final volume of 200 µL/well of CCM medium, added of the
test molecule at 50 µM. DMSO 0.5% (v/v) was used as the control. The plates were incubated in the dark under stationary conditions for 6 days at 25 °C. Aflatoxin accumulation was monitored directly in the culture medium by fluorescence emission determination using a TECAN SpectraFluor Plus microplate reader (Männedorf, Switzerland; λ<sub>ex</sub> = 360 nm; λ<sub>em</sub> = 465 nm; manual gain = 83; lag time = 0 µs; number of flashes = 3; integration time = 200 µs). Samples were inoculated in quadruplicate.

3.10. A. flavus Radial Growth Determination

A. flavus radial growth on a fermentable and a non-fermentable carbon source was performed in YES-agar medium supplemented with 2% glucose (w/v) and 2% ethanol (v/v) respectively, and amended with mHtcum or antimycin A at the concentration of 50 µM. Three single spots (5 µL of a 10<sup>7</sup> conidia/mL suspension each) of the aflatoxigenic strain were equidistantly inoculated in Petri dishes (9 cm Ø), while plates were reviewed in triplicate and incubated for 4 days at 25 °C, and the mycelium growth was evaluated daily by measuring colonies reverse along two orthogonal diameters. The radial growth rate was expressed as percentage inhibition with respect to the control (±SD).

3.11. Sclerotia Biogenesis Assay

A volume of 5 µL from a 10<sup>5</sup> spores/mL suspension of A. flavus aflatoxigenic strain was point inoculated centrally on Czapek Agar plates [1.5% (w/v) agar, 0.1% (w/v) di-potassium hydrogenphosphate, 0.05% (w/v) magnesium sulfate heptahydrate, 0.05% (w/v) potassium chloride, 0.3% (w/v) sodium nitrate, 3% (w/v) sucrose] added with mHtcum or antimycin A at the concentration of 50 µM. Plates were incubated up to 10 days to obtain sclerotia production; then, the surface of colonies was scratched, and sclerotia were manually recovered, ethanol-washed to remove conidia, dried up to 3 days at 60 °C and weighted. Data were recorded as dry weight (mg) and converted into sclerotia production percentage inhibition with respect to the control (±SD). Plates were inoculated in four replicates.

3.12. RNA Extraction and Gene Expression Analysis

Total RNA was extracted from 200 mg of mycelium sampled from 96-h-old microplate CCM cultures treated with mHtcum and antimycin A 50 µM, using a TRIzol® Kit (Sigma-Aldrich, Saint Louis, MO, USA) according to the manufacturer’s instructions. Reverse-transcription of 2 µg of total RNA was conducted using the Maxima First-Strand cDNA Synthesis Kit for qRT-PCR with dsDNase (Thermo Fisher Scientific, Waltham, MA, USA), following the manufacturer instructions. The complementary DNA samples were used as templates of qPCR reactions conducted with ABI 7300 instrumentation (Thermo Fisher Scientific, Waltham, MA, USA) and iTaq™ Universal SYBR® Green Supermix (BioRad, Hercules, CA, USA). Primer sequences, already described in Degola et al. [33], are reported in Table 1. Target genes expression was normalized to the expression of tub1 gene. The ∆CT was calculated as CT target gene − CT internal standard; the expression level variations were then expressed as 2<sup>−∆∆CT</sup> (with ∆∆CT = CT treatment − CT control). Three biological and three technical replicates per condition were performed. Amplification conditions were: 50 °C for 2 min and 95 °C for 10 min; 40 cycles at 95 °C for 15 s and 60 °C for 1 min; 95 °C for 15 s, 60 °C for 1 min, 95 °C for 15 s, and 60 °C for 15 s. The significance of differences between relative expression ratios of treated and control samples was determined using the Mann-Whitney test (p ≤ 0.05).
Table 1. Oligonucleotides sequence and target genes used in this study.

| Oligo Name     | Sequence (5′→3′)          | Target Gene |
|----------------|---------------------------|-------------|
| tubFw-RT       | TACCATGGACGCCGTCCG         | Tub1        |
| tubRev-RT      | GACGGACAACATCGACAAC       |             |
| OmtB F         | GCCCTGACATGGAAACCATC      |             |
| OmtBRev-RT     | TCCACTGCTCAATCGCATG       |             |
| Adh1 FwRT      | CTAACACCAGGACAGATGAG      | Adh1        |
| Adh1 RevRT     | TCCCTCGTGTCCACCTAC        |             |
| Nor1 F         | ACGGATCAGTCGACGCCAC       |             |
| AflD-RevRT     | ACGGTGCCTTGGGACGTTG       | AflD        |
| AflJ-gF        | GAACGCTGATTGCCAATGCC      | AflS        |
| AflS RevRT     | GATTCATCCAGGAGGATAAC      |             |
| VeA FwRT       | CGAGACGGAAAGCCTCCGT       | VeA         |
| VeA RevRT      | TGGAGGATCGACTGGACGA       |             |

3.13. Statistical Analysis

For statistical analyses, one-way analysis of variance (ANOVA) was used in the Past 3.x software. Results of mycelial growth, aflatoxin accumulation, and sclerotia production were analyzed by Tukey’s test; differences were considered significant at $p \leq 0.05$.

4. Conclusions

Fungi secondary metabolism and, in particular, mycotoxins biosynthetic pathways require a well-defined regulation of oxidative intracellular balance triggering the redox homeostasis needed for both fungal development and differentiation [52]. Because of this pivotal role, destabilization of oxidative-linked systems, such as mitochondrial electron flow, can be an effective way to control the fungal production of aflatoxins. The data presented in this paper suggest that the toxin inhibition induced by mHtcum is linked to molecular switches regulating the carbon source utilization and the energy production in the cell. The interaction between the compound and bc1 cytochrome in the active site of mitochondrial complex III, proposed in consideration of the results obtained in the yeast model system, has been supported by a docking analysis validation. On the other hand, since in A. flavus aflatoxin biosynthesis and sclerotia biogenesis, that are both coordinately inhibited by mHtcum treatment, do not equally respond to a specific inhibitor of complex III, the existence of multiple (at least two) targets for thiosemicarbazone should be presumed. Thus, we propose that mHtcum might operate on a regulatory step that is located upstream of the bifurcation of aflatoxin biosynthesis and sclerotia development pathways. However the overall regulatory machinery might be more sophisticated since it should be considered that, as a mycelia forming fungus, Aspergillus could go through differentiation/specialization processes, resulting in a task division (some cells engage in energy metabolism, whereas other do so in the secondary metabolites production).

Supplementary Materials: The following are available online. Figure S1: Effect of mHtcum on yeast cells growth on different non-fermentable carbon sources. In the disc diffusion assay, ethanol was replaced by acetate or glycerol as carbon source in the YP medium.

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**Sample Availability:** Samples of the compounds are available from the authors.