Search for the Active Ingredients from a 2-Aminothiazole DMSO Stock Solution with Antimalarial Activity

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Chemical decomposition of DMSO stock solutions is a common incident that can mislead medicinal screening campaigns. Here, we share our case study of 2-aminothiazole 1, originating from an antimalarial class that undergoes chemical decomposition in DMSO at room temperature. As previously measured biological activities observed against *Plasmodium falciparum* NF54 and for the target enzyme PfIspE were not reproducible for a fresh batch, we tackled the challenge to understand where the activity originated from. Solvent- and temperature-dependent studies using HRMS and NMR spectroscopy to monitor the decomposition led to the isolation and in vitro evaluation of several fractions against PfIspE. After four days of decomposition, we successfully isolated the oxygenated and dimerised compounds using SFC purification and correlated the observed activities to them. Due to the unstable nature of the two isolates, it is likely that they undergo further decomposition contributing to the overall instability of the compound.

In the search for novel antimalarial compounds targeting the kinase IspE of the 2-C-methyl-D-erythritol 4-phosphate (MEP) pathway, we identified a new 2-aminothiazole class via an enzymatic high-throughput screening (HTS) campaign (unpublished results from the consortium). The MEP pathway is a key biosynthetic route for the production of universal isoprenoid precursors.[1] The HTS yielded compound 1 as a hit that was follow-up due to its promising screening profile (Table 1), which was supported by some previously reported 2,4-substituted thiazoles with antiplasmodial activity.[2,3]

For a newly synthesised batch and the corresponding freshly prepared stock solution of compound 1, no activity against *Plasmodium falciparum* (Pf) IspE was detected and cell-based activity against the plasmodial strain PNF54 resulted in an 17-fold reduction in activity (Table 1). Previous samples originated from older plates stored in DMSO, which had undergone several thawing cycles from –20°C storage. First evidence of the decomposition was visually observed due to a change in colour of the compound’s stock solution from clear to dark. However, this colour change would only be obvious to someone familiar with the original colour of the parent compound. If such plates are sent to collaboration partners responsible for biological assays, as is often the case in medicinal chemistry projects, such alterations may not necessarily be observed or questioned. DMSO is a widely used solvent due to its amphiphilic nature, enabling higher solubility of many compounds and high viscosity improving the reproducibility in pipetting. Nevertheless, stability issues of chemical compounds kept as stock solutions in DMSO are also acknowledged and spontaneous reactions, such as oxidation, cyclisation and hydrolysis, in stock solutions may affect the biological activity originated from. Solvent- and temperature-dependent studies using HRMS and NMR spectroscopy to monitor the decomposition led to the isolation and in vitro evaluation of several fractions against PfIspE. After four days of decomposition, we successfully isolated the oxygenated and dimerised compounds using SFC purification and correlated the observed activities to them. Due to the unstable nature of the two isolates, it is likely that they undergo further decomposition contributing to the overall instability of the compound.

**Table 1.** Starting point for the non-reproducible results of compound 1.

| Old batch (decomposed) | 8.0 ± 2.8 | 0.43 ± 0.01 |
| Fresh batch | > 500 | 7.3 ± 0.8 |

[a] Contribution for this article is available on the WWW under https://doi.org/10.1002/cmdc.202100067
activity.\(^{4-6}\) Although reactivity of 2-aminothiazoles has not been directly ascribed to DMSO, we were concerned about it. 2-Aminothiazoles are frequently-hitting fragments in biophysical binding assays, as the so-called promiscuous 2-aminothiazoles (PrATs), and some sub-categories are classified as Pan-Assay Interference Compounds (PAINS).\(^{19-21}\) The observed activity on target in vitro and in whole-cell systems prompted us to elucidate the structural changes of compound 1 in DMSO that accounted for the increased inhibitory activity.

Firstly, we performed a temperature-dependent decomposition study of compound 1 at 10 mM concentration in DMSO at room temperature (RT), \(+4\) °C and \(-20\) °C (SI, Section 2.3). After seven days, 64% of the sample stored at RT had decomposed (SI, Table S1). Besides visual inspection of the colour changes of the test sample (see Table 2), the decomposition was monitored by HRMS and NMR spectroscopy. The sample that was incubated at \(-20\) °C underwent only minimal decomposition, whereas, the sample kept at RT was fully decomposed after two weeks (SI, Figure S10). Furthermore, all three samples were tested for their inhibitory activity against PfISP\(_E\) after the incubation period. Only the samples incubated at RT and \(+4\) °C showed measurable inhibitory activity in the PfISP\(_E\) assay, which was not observable for the freshly prepared compound 1 (Table 2). This fact correlated with the degree of chemical transformation of compound 1 (Table S1). In the IspE assay, the activity of the target enzyme is coupled to the oxidation of NADH (which can be followed spectrophotometrically at 340 nm) via a cascade of the auxiliary enzymes pyruvate kinase and lactate dehydrogenase (PK/LDH).\(^{22,23}\) Therefore, we next confirmed, whether the effects observed in the IspE assay are due to inhibition of the target enzyme or of the auxiliary enzymes. Three samples of compound 1 were tested in PK/LDH assay (Table 2). The resulting IspE values of 34 \(\mu\)M and 45 \(\mu\)M for compound 1 incubated at RT and \(+4\) °C, respectively, are only about three times higher than the IC\(_{50}\) values determined in the PfISP\(_E\) assay (12 \(\mu\)M and 16 \(\mu\)M, respectively). From these results, it was impossible to evaluate the influence of inhibition of auxiliary enzymes on the IC\(_{50}\) values determined in the PfISP\(_E\) assay.

Another approach to address this issue is to perform the IspE assay with one or several orthologs of PfISP\(_E\). The inhibitory potency of samples incubated at different temperatures of compound 1 against Escherichia coli (Ec)IspE are shown in the Table 2. The IspE assay setup was identical for both Pf and EcIspE orthologs except for the target enzyme used. One of the setup requirements was that the enzymatic activity of the auxiliary enzymes PK/LDH exceeded the activity of the target enzyme IspE not less than a factor of ten.\(^{110}\) We investigated, if the inhibition of the auxiliary enzymes may substantially influence the IC\(_{50}\) values obtained from the coupled IspE assay. The IC\(_{50}\) values for the active batches (“RT” and \(+4\) °C) of compound 1 observed in the EcIspE assay are eight- and four-fold less active in comparison to those observed in the PfISP\(_E\) assay (Table 2). The fact that the inhibition of auxiliary enzymes in the EcIspE assay did not give rise to equally low IC\(_{50}\) values in comparison to those from PfISP\(_E\) assay means that the inhibition of auxiliary enzymes influenced only marginally the IC\(_{50}\) values measured in the PfISP\(_E\) assay.

Secondly, we analysed if spontaneous chemical transformation of compound 1 may depend on the solvent used for the preparation of its stock solutions. We incubated 10 mM stock preparations of compound 1 in DMSO, acetonitrile (ACN) and methanol (MeOH) at RT. Aliquots were taken during 16 days of incubation and tested against the enzyme PfISP\(_E\) (Table 3). Interestingly, no PfISP\(_E\) inhibitory activity for the ACN or MeOH aliquots was observed, meaning that no chemical transformation of compound 1 took place in ACN or MeOH, as supported by the HRMS and NMR data (SI, Section 2.4). For further decomposition incubation of compound 1, only DMSO was used as solvent.

The HRMS chromatographic profile of the analytical sample of compound 1 after 16 days of incubation at RT showed decomposition into several peaks, as seen in the chromatographic profiles of the samples previously used for the biological testing (SI, Figure S13). For further analysis, we repeated the incubation in DMSO on a larger scale for two weeks and fractionated the mixture by semipreparative HPLC into 29 fractions. The inhibitory activities of the fractions were determined against PfISP\(_E\) (SI, Table S2). In the preparative sample, a peak (m/z: 426.12607) was detected and interestingly, 

| Table 2. Temperature-dependent decomposition of compound 1 in DMSO. |
|-------------------|-----------|-----------|-----------|-------------|
| Activity measured after 3 months | RT\[a\] | \(+4\) °C\[b\] | \(-20\) °C\[b\] | 1 - Old sample\[c\] |
| PfISP\(_E\) IC\(_{50}\) (\(\mu\)M)\[d\] | 12 ± 4 | 16 ± 7 | > 500 | 10 ± 3 |
| PK/LDH IC\(_{50}\) (\(\mu\)M)\[d\] | 34 ± 4 | 45 ± 6 | n.d. | 40 ± 4 |
| EcIspE IC\(_{50}\) (\(\mu\)M)\[d\] | 101 ± 14 | 71 ± 10 | > 500 | 32 ± 6 |
| Degradation | 100% | 79% | 18% | n/A |

[a] Errors given as formal standard error. [b] HRMS chromatograms measured shortly before the assay are given in SI, Figure S7. [c] The control values for the old decomposed sample run at the same time: n.d.: not determined, N/A: not applicable, PK/LDH: pyruvate kinase and lactate dehydrogenase, Pf: Plasmodium falciparum, Ec: Escherichia coli, RT: room temperature.

| Table 3. Time-dependent decomposition of compound 1 at room temperature. |
|-------------------|-----------|-----------|-----------|-------------|
| Days of incubation | DMSO | PfISP\(_E\) IC\(_{50}\) (\(\mu\)M)\[a\] | PK/LDH IC\(_{50}\) (\(\mu\)M)\[a\] | ACN | MeOH | DMSO |
| 0 | > 500 | > 500 | > 500 | n.d. |
| 5 | 19 ± 7 | > 500 | > 500 | n.d. |
| 7 | 24 ± 9 | > 500 | > 500 | 48 ± 16 |
| 9 | 10 ± 4 | > 500 | > 500 | 57 ± 22 |
| 12 | 1 ± 1 | > 500 | > 500 | 2 ± 1 |
| 14 | 10 ± 4 | > 500 | > 500 | 50 ± 11 |
| 16 | 11 ± 4 | > 500 | > 500 | 86 ± 17 |
| Solvent Blank | > 500 | > 500 | > 500 | n.d. |

[a] Errors given as formal standard error. PK/LDH: pyruvate kinase and lactate dehydrogenase, Pf: Plasmodium falciparum. n.d.: not determined.
the sample incubated in DMSO-d₄ included a peak at 4.86 min with the likely addition of the deuterated methyl groups to the mass (m/z: 432.16354), which initially attracted our attention due to a possible reaction with DMSO itself. However, this peak was isolated from the DMSO sample as fraction 4 in the semipreparative HPLC and it did not show any inhibitory activity against PfISP (SI, Table S2). Out of the isolated fractions, most of the active degradation products are very non-polar and poorly separable by rp-HPLC, showing dimerised masses as well as a mass of 380.10302/380.2 Da. The previous HRMS studies revealed that these signals appeared already after a couple of days of incubation at RT. In particular, we could observe that two peaks (m/z: 380.10257 and m/z: 365.11542) appeared after 15 hours and the latter becomes more prominent after four and a half days (SI, Figure S22).

Thus, we performed another round of large-scale decomposition in DMSO at RT for five days and optimised the semipreparative HPLC conditions (SI, Table S3). A second purification step with supercritical fluid chromatography (SFC) (SI, Section 2.6) finally yielded two main active degradation products in sufficient amount for NMR analysis: decomposition product (DP1) with the m/z value of 380.10376 [M+H]+ or [M]+, corresponding to the sum formula of C₁₇H₁₇F₃N₆O₁₈ and respectively C₁₇H₁₇F₃N₆O₁₈ (PfISP IC₅₀: 199 ± 26 μM) and DP2 with the m/z of 365.11542 [M+2H]+; corresponding to the sum formula of C₁₇H₁₇F₃N₆S₂⁺ (PfISP IC₅₀: 59 ± 4 μM) (Table 4, Figure 1).

DP1 has a very distinct orange colour, supporting the formation of the formed oxygenated thiazolone core (UVmax = 196, 223 and 457 nm), (SI, Figure S32). The addition of the oxygen atom was evident based on the MS data and the exact position was confirmed by the disappearance of the characteristic -CH proton signal of the thiazole 5-position at 6.54 ppm in the ¹H NMR spectrum (SI, Table S4, Figure S24–S25) supported by a change in chemical shift of the neighbouring carbon atom in 4-position from 141.3 ppm to 172.5 ppm. In the parent compound 1, HMBC correlation of the thiazole carbon in 4-position (141.3 ppm) with the proton 5-position (6.52 ppm) is clear, whereas in the DP1, the thiazole carbon in 4-position (172.5 ppm) correlates with the nearest methyl of the pyrrole (2.83 ppm) (SI, Figure S28–S29). Detection by UV afforded a single peak, which, however, underwent isomerisation into a more polar derivative over time. In fact, the long-term stability of DP1 became questionable as different stabilities were observed in acetone, MeOH, ACN, DMSO and CDCl₃ (SI, Figure S33–S37). Chloroform induces addition of chlorine as evidenced by the isotopic pattern, (m/z: 462.0 and 464.0 for [²Cl] and [³Cl], respectively, Figure S35). The compound was most stable in acetone, in which we recorded NMR spectra and identified a mixture of compounds, as predicted based on the noticeable shift in the retention time in LCMS. The bond between the imine of the thiazolone core and the phenyl ring can rotate, as indicated in Figure 1, and thus, it is likely that DP1 exists as either E or Z isomer. We calculated the energies of the lowest-energy conformations, pointing towards an E configuration of the isolated DP1 (+29.4 kcal/mol). Additionally, the appearance of a broad singlet at 5.84 ppm for -NH and the changes in the pyrrole shifts supports the formation of a charged pyrrolium species that can also exist either as E or Z isomer (+66.1 vs +62.8 kcal/mol). Considering the overall reactivity of the 2,4-thiazole substitution, tautomisation into a more stable conjugated form can occur due to the slightly acidic and dipolar nature of DMSO. This would mean that we measure the charged form in the HRMS as 380.10376 [M]+. Due to the overall stability of DP1, further experiments to confirm the exact isomeric mixtures are cumbersome and were not pursued. Nevertheless, the oxygenated DP1 in its isomeric mixture from the SFC separation showed moderate, yet selective, inhibitory activity against PfISP (IC₅₀ = 199 ± 26 μM) without PK/LDH inhibition (IC₅₀ = > 500 μM).

| Table 4. | The biological data of the SFC-separated compounds and analytical samples over four days. |
| Days of incubation | IC₅₀ (μM) (PfISP) | EclispE | PK/LDH | PNF54 |
|---------------------|-----------------|--------|--------|-------|
| 0                   | >500            | >500   | n.d.   | n.d.  |
| 1                   | >500            | >500   | n.d.   | n.d.  |
| 2                   | >500            | >500   | 3.5 ± 0.3 | n.d.   |
| 3                   | 486 ± 20        | 397±1  | n.d.   | n.d.  |
| 4                   | 99 ± 23         | 75 ± 16| 95 ± 21 | 2.7 ± 0.3 |
| DP1 “380”           | 199 ± 26        | n.d.   | >500   | n.i. (>20) |
| DP2 “365”           | 59 ± 4          | 37 ± 4 | 2.1 ± 0.2 | n.d.   |

[a] All compounds were dissolved in methanol shortly before the assay. [b] Freshly dissolved compound 1 (PfISP IC₅₀ = >500 μM). [c] Error given as formal standard error. n.d.: not determined, n.i.: no inhibition at the highest concentration tested, PK/LDH: pyruvate kinase and lactate dehydrogenase, Pt: Plasmodium falciparum, Ec: Escherichia coli, DP: decomposition product.

Figure 1. Characterised compounds isolated from the decomposition mixture. The shown chromatographic traces are base peak chromatograms generated with supercritical fluid chromatography.
but showed no inhibition in the Pf cell-based assay at the highest tested concentration ([IC₅₀ > 20 μM]). Importantly, the cellular assay occurs over a longer time (72 hours) than the enzymatic assay (30 min), which may influence the overall stability of the compound. In addition, we monitored the HRMS chromatographic profile of freshly dissolved compound 1 in the enzyme assay buffer (Tris-HCl, pH 7.6) with and without 5 % DMSO for 30 min (SI, Figure S16–17). The presence of DMSO seems to influence the formation of DP1 (m/z: 380.10260 and 380.10238) and DP2 (m/z: 365.11554), even within the short time window of 30 min.

Structure elucidation by NMR spectroscopy showed that the mass of DP2 fits in fact to a dimer of 1 (Figure 1). We observed most often the doubly protonated species with m/z of 365.11542 [M + 2H]²⁺ more intensely than the charged mass [M]⁺: 729.22473. With aid of the SFC, we found out that an isomer of DP2 starts to form over the four days of incubation, as seen by the appearance of the minor peak, as highlighted in the green box in Figure 1. NMR measurements revealed its non-symmetry as one of the characteristic –CH peaks corresponding to the thiazole position 5 disappeared and a new peak appeared at 3.96 ppm, integrating for –CH₃ (SI, Table S5). This led us to question how the dimerisation would occur. Based on the NMR and MS data, we propose that an N–N bond formation occurs between the linker amines, as represented in the proposed reaction (SI, Scheme S1). N–N bonds are generally unstable, however, rather commonly occur in biological complexes.[11] Due to the slightly acidic nature of DMSO, it is likely that the conjugated pyrrolium species is present and the imine N, as in tautomer of compound 1, is attacked by the nucleophilic linker amine forming the N–N bond. The charged pyrrolium may also undergo isomerisation between E and Z isomers as observed in the SFC conditions, (SI, Figure S38). Z being slightly more stable than E (+144.7 vs +148.5 kcal/mol), hampering the exact assignment of the pyrrolium NMR peaks.

For DP2, we could reach a similar enzymatic activity profile as for the initial decomposed starting point (PfSpE IC₅₀ = 59 ± 4 μM), although also inhibiting the auxiliary enzymes (PK/LDH from a single measurement) that was not reproducible later for a freshly dissolved batch (PfSpE IC₅₀ > 500 μM), reinforcing the need for careful handling of any related compounds.

In conclusion, we analysed the collected data to understand what causes the antimalarial activity that evolved from compound 1 stored in DMSO. The PfSpE activity can be partly ascribed to dimerised DP2, although it also inhibits PK/LDH. On the other hand, the oxygenated DP1 may also play a role when in its active isomeric form. The isolated degradation products themselves are unstable. They undergo further degradation, leading to a mixture that may also account for the observed activities. Additionally, our study underlines the importance of appropriate storage conditions of DMSO stock solutions of 2-aminothiazoles. Based on the temperature-dependent studies, we confirmed that decomposition hardly occurs at ~20 °C over two months in DMSO. For the future, blocking the 5 position of the thiazole ring with a fluorine atom could be a feasible strategy to reduce the reactivity.[12] However, to avoid decomposition, working with such a class requires preparation of fresh stock solutions prior to biological assays. Furthermore, multiple freeze-thaw cycles should be avoided and alternative solvents should be considered. With this communication, we wish to remind the medicinal-chemistry community again that what is in the test well, might not be the compound one thought.
Experimental Section
Details for chemical syntheses, analytical and biological methods together with characterisation data are described in the Supporting Information.

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Conflict of Interest
The authors declare no conflict of interest.

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