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Synthesis, Mechanism Elucidation and Biological Insights of Tellurium(IV)-Containing Heterocycles

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Abstract: Inspired by the synthetic and biological potential of organotellurium substances, we describe here the synthesis and characterization of a series of five- and six-membered ring organotelluranes containing a Te–O bond. Theoretical calculations elucidated the mechanism for the oxidation-cyclization processes involved in the formation of the heterocycles, consistent with chlorine transfer to hydroxy telluride, followed by cyclization step with simultaneous formation of the new Te–O bond and the OH group deprotonation. Moreover, theoretical calculations also indicated *anti*-diastereoisomers as major products for two-chirality center-containing compounds. Antileishmanial assays against *Leishmania amazonensis* promastigotes disclosed 1,2 λ^4 -oxatellurane **LQ50** (IC₅₀ = 4.1 $\pm$ 1.0; SI = 12), 1,2 λ^4 -oxatellurolane **LQ04** (IC₅₀ = 7.0 $\pm$ 1.3; SI = 7) and 1,2 λ^4 -benzoxatellurolane **LQ56** (IC₅₀ = 5.7 $\pm$ 0.3; SI = 6) as more powerful and more selective compounds, up to four times more active than the reference. Stability study supported by ¹²⁵Te NMR analyses showed that these heterocycles do not suffer structural modifications in aqueous-organic media and up to 65 °C.

Introduction

Tellurium is the only natural chalcogen with no apparent role in vital biological processes. Tellurium toxicity for humans still remains poorly understood, as its biological role is probably less known than metals like osmium and ruthenium.^[1] Some of the few reports address the use of tellurium for the synthesis of amino acids and proteins in fungi and bacteria.^[2] Despite the presence of tellurium in living organisms occurring only in exceptional cases, synthetic tellurium-containing compounds have been applied in biological systems^[3] and reported as promising antioxidant agents,^[4] immunomodulator,^[5] antimicrobial,^[6] anti-

inflammatory,^[7] antitumoral^[8] and enzymatic inhibitors.^[9] Additionally, this element has been used for several applications in organic and inorganic chemistry, like materials and medicinal fields.^[1,10]

As well established, tetra- and hexa-valent tellurium-containing compounds (Te(IV) and Te(VI)) are exceptions to the Lewis octet rule,^[11] being classified as hypervalent compounds.^[12] Such hypervalent derivatives present great potential for biological purposes.^[13] For decades, numerous studies revealed that the inorganic tellurane **AS101** (Figure 1) is a potent inhibitor of the cysteine proteases such as papain and cathepsin B.^[9a,14] From then on, several organic and inorganic hypervalent tellurium compounds were reported as successful enzyme inhibitors, including cysteine,^[6g, 9b-c, 15] and threonine proteases (20S proteasome),^[9h] and tyrosine phosphatases (PTP1B and YopH),^[9f] also indicated as promising antioxidants,^[4d] antimicrobial^[6c,e, 16] and antitumor agents.^[8f]

Another important example of a biologically active tellurium-containing compound is the organotellurane **RT-01** (Figure 1), which presented *in vivo* antiprotozoal activity against *Leishmania amazonensis*, leading to significant decrease of the parasitic load compared to that of the reference drug.^[17] Another organotellurane, **RF07** (Figure 1), was described as a powerful leishmanicidal agent for *Leishmania chagasi* amastigotes with selectivity index (SI = CC₅₀/IC₅₀) of 10, relating the specificity of the activity in comparison to the cytotoxicity of the evaluated compound.^[18] This compound was previously described as a cysteine protease cathepsin B inhibitor,^[9b] and its antileishmanial activity has been associated with inhibition of this enzyme in *L. chagasi* protozoa. It is worth mentioning that the inorganic tellurane **AS101** (Figure 1) also presented antileishmanial activity, with IC₅₀ = 26.9 μ mol L⁻¹ for *Leishmania donovani* promastigotes.^[14f]

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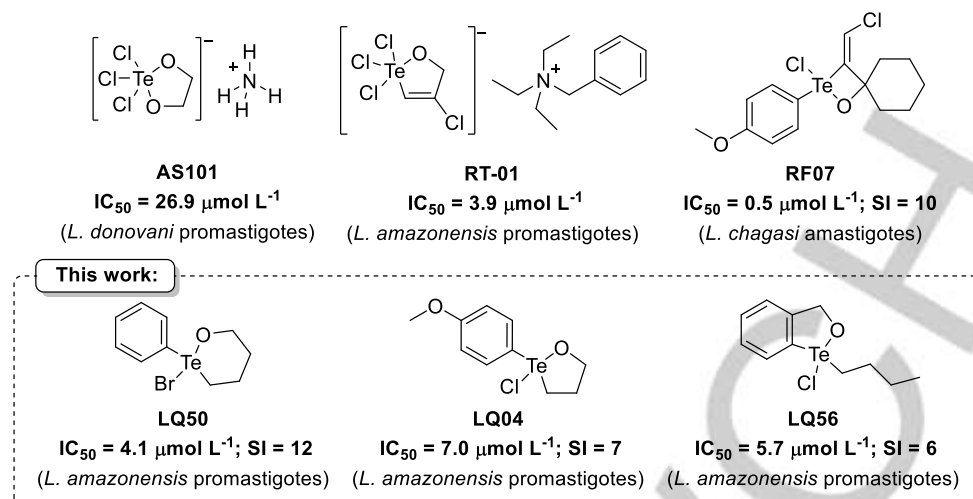


Figure 1. Chemical structures and antileishmanial activities of hypervalent tellurium compounds.

Hypervalent five- and six-membered Te–O bond-containing heterocycles have already been described in the literature;^[19] however, their biological properties remain poorly explored. Then, inspired by the potential of hypervalent tellurium compounds, in the current work we discuss: i) the synthesis of a series of five- (1,2 λ^4 -oxatellurolanes and 1,2 λ^4 -benzoxatellurolanes) and six- (1,2 λ^4 -oxatelluranes) membered ring organotelluranes; ii) calculations using Density Functional Theory (DFT), performed to identify the most favorable mechanism and to rationalize the diastereoselectivity observed for two-chirality center-containing compounds; iii) the *in vitro* antileishmanial potential of all hypervalent Te(IV)-containing compounds, evaluated against *L. amazonensis* promastigotes; and iv) stability assays supported by ¹²⁵Te NMR (thermal stability and chemical stability in aqueous-organic media) for some representative hypervalent Te(IV)-containing compounds.

Results and Discussion

Synthesis of Tellurium(IV)-Containing Heterocycles

The synthetic routes to access 1,2 λ^4 -oxatellurolanes and 1,2 λ^4 -oxatelluranes were performed as shown in Scheme 1. Bromobenzene (**1a**) or 4-substituted aryl bromides (**1b–e**) were reacted with elemental magnesium, followed by the addition of elemental tellurium and later air oxidation, leading to diaryl ditellurides **2a–e** in 27–65% isolated yields. The Te–Te bonds from ditellurides **2a–e** were cleaved employing sodium borohydride and their respective tellurolates were captured by 3-chloropropan-1-ol to produce hydroxy tellurides **3a–e** or by 4-bromobutan-1-ol leading to hydroxy telluride **3f**. Finally, one-pot oxidation-cyclization sequential reactions of **3a–f** employing *N*-chlorosuccinimide (NCS) or *N*-bromosuccinimide (NBS) led to corresponding 1,2 λ^4 -oxatellurolanes (**LQ02–04**, **33–35** and **48**) in 2–79% isolated yields and 1,2 λ^4 -oxatelluranes (**LQ49–50**) in 5–15% isolated yields (Scheme 1).

5-Methyl- (**LQ51** and **LQ52**) and 5-phenyl-substituted 1,2 λ^4 -oxatellurolanes (**LQ53**) were prepared by a 1,4-addition of phenyl tellurolate (from Te–Te bond cleavage of **2a** with NaBH₄) to but-3-en-2-one or 1-phenylprop-2-en-1-one, followed by *in situ* carbonyl reduction with NaBH₄. In the sequence, NCS or NBS mediated oxidation-cyclization sequential reactions of hydroxy

tellurides **3g–h** led to 1,2 λ^4 -oxatellurolanes **LQ51–53** in 30–79% isolated yields (Scheme 2).

Hydrolysis of the acetal moiety of **3e** led the hydroxy telluro ketone **3i** (90% isolated yield), which was applied as an intermediate for the hydroxy telluro oxime **3j** (82% isolated yield). One-pot sequential oxidation-cyclization sequential using NCS led to the 1,2 λ^4 -oxatellurolanes **LQ54** and **LQ55** in 30 and 24% isolated yields, respectively (Scheme 3).

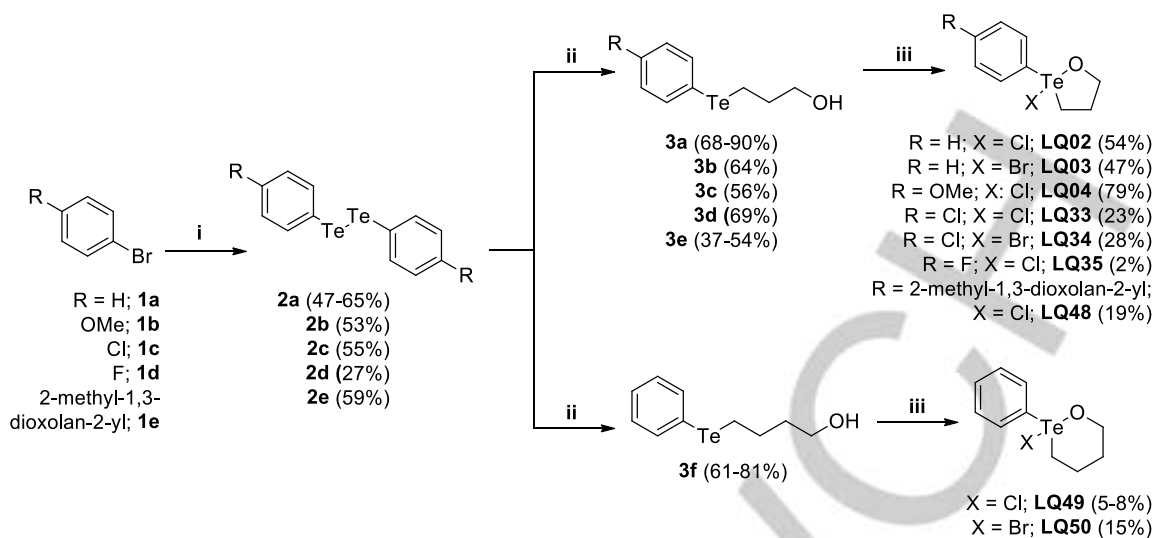
2,5-Dihydro-1,2 λ^4 -benzoxatellurolanes **LQ56** and **LQ57** were prepared in 30 and 24% isolated yield, respectively, employing the same oxidation-cyclization approach with NBS or NCS from benzyl hydroxy telluride **3k** (Scheme 4).

All hypervalent Te-containing heterocycles (**LQ02–04**, **LQ33–35** and **LQ48–57**) were purified by precipitation in H₂O/Dimethyl sulfoxide (10:1 v/v) followed by second precipitation in hexane/CH₂Cl₂ (10:1 v/v), and were obtained as odorless white solids with good solubility in organic solvents such as dichloromethane, acetone, tetrahydrofuran, and dimethyl sulfoxide. Eleven chlorinated and five brominated compounds (thirteen unpublished) were obtained in 2–79% isolated yields in the oxidation-cyclization step and overall yields ranging from 0.4% (**LQ35**) to 40% (**LQ52**). Structural characterizations were performed by melting point determination, ¹H Nuclear Magnetic Resonance spectroscopy (NMR), ¹³C NMR, ¹²⁵Te NMR, Fourier-transform infrared (FTIR) spectroscopy, and single-crystal X-ray diffraction (if applicable).

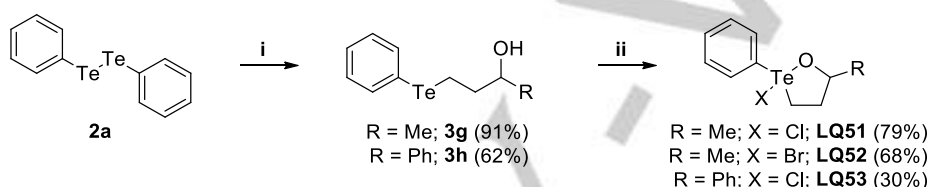
Structural Characterization by NMR

Rotational restriction and a slightly bent conformation of the five-membered rings in 1,2 λ^4 -oxatellurolane forced the geminal methylene hydrogens to assume pseudo-axial and pseudo-equatorial positions. Furthermore, tetrasubstituted Te-atom is a chirality center and the methylene hydrogens in the heterocycles are diastereotopic. These structural features are easily observed in three-dimensional projections (Figure 2a, b) and NMR data of representative 1,2 λ^4 -oxatellurolanes **LQ02** (Figure 2c) and **LQ03** (Figure 2d). Geminal methylene hydrogens presented distinct signals that confirm the heterocyclization as can be seen in correlation maps in Figure 2c and 2d. The multiplicity of hydrogens at position 1 revealed the geminal coupling between them ($J = 10.0$ Hz) and vicinal couplings with the hydrogens at position 2 ($J = 5.8, 4.0$ and 3.4 Hz). The same pattern of multiplicity was observed for hydrogens at position 3, with geminal

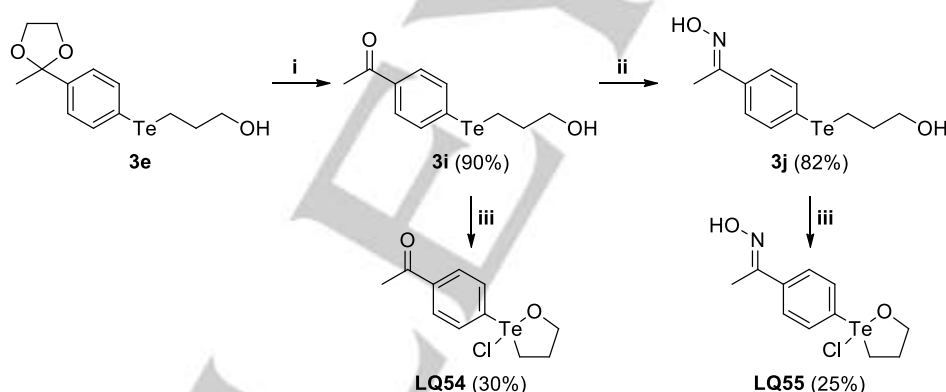
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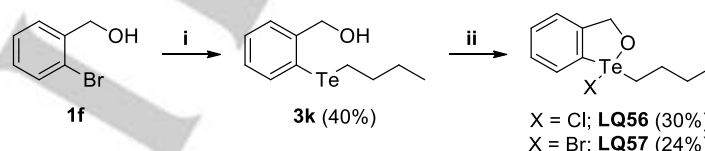
Scheme 1. Synthesis of 1,2λ⁴-oxatellurolanes **LQ02-04**, **33-35** and **48** and 1,2λ⁴-oxatelluranes **LQ49-50**. Reagents and Conditions: i. 1. Mg⁰, I₂, tetrahydrofuran, r.t., 2 h; 2. Te⁰, 2 h; 3. O₂, 15 min. ii. 3-chloropropan-1-ol or 4-bromobutan-1-ol, NaBH₄, H₂O/EtOH, 0 °C-r.t., 1 h. iii. NCS or NBS, CH₂Cl₂, 0 °C-r.t., 30 min.



Scheme 2. Synthesis of 1,2λ⁴-oxatellurolanes **LQ51-53**. Reagents and Conditions: i. 1. NaBH₄, H₂O/EtOH, 0 °C, 5 min. 2. α,β-unsaturated ketone, 0 °C-r.t., 1 h; 3. NaBH₄, 0 °C-r.t., 15 min. ii. NCS or NBS, CH₂Cl₂, 0 °C-r.t., 30 min.



Scheme 3. 2-Methyl-1,3-dioxolan-2-yl group derivatization for the synthesis of 1,2λ⁴-oxatellurolanes **LQ54** and **LQ55**. Reagents and Conditions: i. *p*-toluenesulfonic acid (pTSA), acetone, reflux, 2 h. ii. (NH₃OH)Cl, AcONa, MeOH, reflux, 2 h. iii. NCS, CH₂Cl₂, 0 °C-r.t., 30 min.



Scheme 4. Synthesis of 1,2λ⁴-benzoxatellurolanes (**LQ56** and **LQ57**). Reagents and Conditions: i. 1. BuLi, THF, -78 °C, 30 min; 2. (BuTe)₂, r.t., 4 h. ii. NCS or NBS, CH₂Cl₂, 0 °C-r.t., 30 min.

(*J* = 12.1 Hz) and vicinal couplings (*J* = 10.6, 8.2, 7.1 and 3.7 Hz). Thereby, the hydrogens on position 2 showed a complex signal composed by their geminal coupling and four more couplings with close *J* values. The most deshielding hydrogens (at 2.14, 3.70 and 4.75 ppm) occupy pseudo-axial positions for this conformation while the most shielding hydrogens (at 1.90, 3.47 and 3.68 ppm) occupy pseudo-equatorial positions. The chemical

shifts, multiplicity, and *J* values for **LQ 02** are detailed in Figure 2e.

¹³C{¹H} NMR spectra comparison of starting material (**3a**) and product (**LQ02**) disclosed significant changes in chemical shifts, mainly for the carbons directly attached to the tellurium atom. Chemical shifts of aliphatic and aromatic carbon atoms (positions

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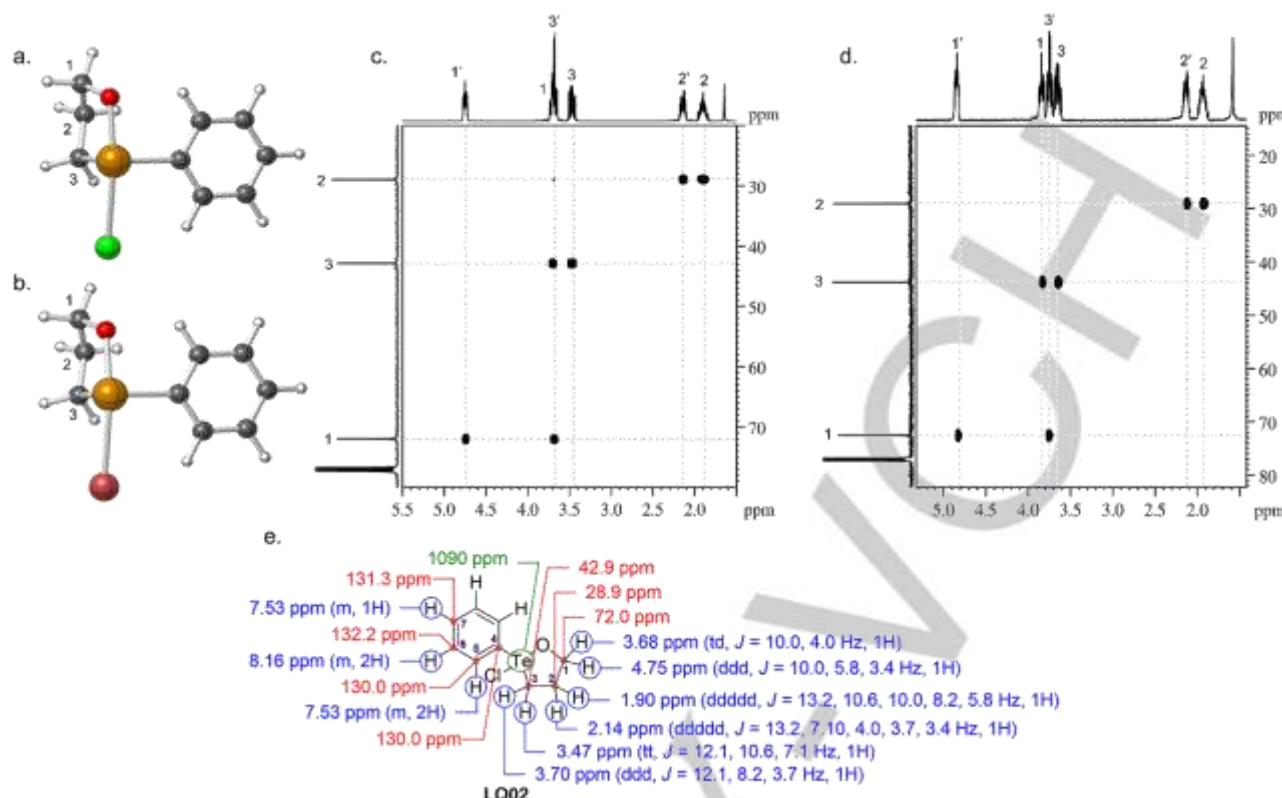


Figure 2. Three-dimensional structure of 1,2λ⁴-oxatellurulane **LQ02** (a) and **LQ03** (b; atom-color: Te-yellow, O-red, Cl-green, Br-dark red, C-gray, H-white.), ¹H-¹³C HSQC correlation maps for **LQ02** (c) and **LQ03** (d) and chemical shift on ¹H (blue), ¹³C (red) and ¹²⁵Te (green) NMR analyses for **LQ02** (e).

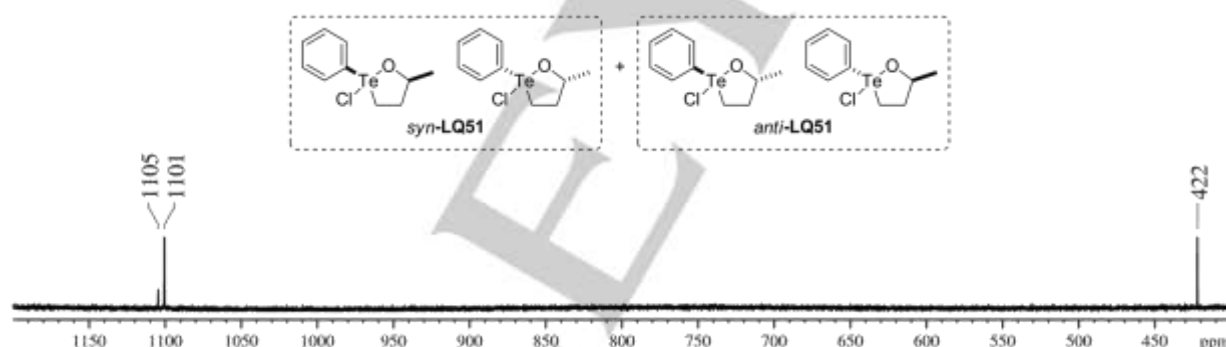


Figure 3. ¹²⁵Te NMR spectrum (126.24 MHz, CDCl₃, PhTeTePh (δ = 422)) of 1,2λ⁴-oxatellurulane **LQ51**.

3 and 4) changed from 5.1 and 109.4 ppm to 43.2 and 128.1 ppm, respectively, after oxidation of hydroxy telluride **3a** to **LQ02** (See Supporting Information). The most significant difference of chemical shift after the oxidation step was observed in ¹²⁵Te NMR analyses, in which the chemical shift varied from 473 ppm (**3a**) to 1090 ppm (**LQ02**).

Formation of diastereoisomers for two-chirality center-containing compounds was confirmed by the presence of two signals in ¹²⁵Te NMR spectra, as exemplified by **LQ51** spectrum in Figure 3. The difference in intensity between those signals demonstrated some degree of diastereoselectivity during the cyclization step. Signals at δ = 1101 and δ = 1105 are in accordance with chemical shifts obtained for the other synthesized tellurium(IV)-containing heterocycles and indicate that one of diastereoisomers (*syn*-**LQ51** or *anti*-**LQ51**) was preferentially formed.

Single-crystal X-ray Crystallography

Single-crystal X-ray diffraction analysis for representative compounds confirmed the cyclization products (Figure 4). Both

LQ02 (Figure 4a) and **LQ51** (Figure 4c) presented two independent molecules, which are enantiomers, in their crystals, and **LQ49** (Figure 4b) and **LQ56** (Figure 4d) are in centrosymmetric space groups and these crystals therefore contain both enantiomers too. Selected bonds and crystallographic data are listed in Tables 1 and 2. The structures revealed that all the compounds adopted a seesaw-molecular geometry about the Te atom, typical for the hypervalent center of Te(IV).^[19c, 20] Hypervalent O–Te–Cl bonding presented average 2.00 and 2.66 Å lengths for Te–O (Te1–O1 and Te2–O2, entries 1 and 2, Table 1) and Te–Cl (Te1–Cl1 and Te2–Cl2, entries 3 and 4) with an average 170° angle (O1–Te1–Cl1 and O2–Te2–Cl2, entries 9 and 10). Te–C bonds at the equatorial positions had an average 2.13 Å length (entries 5–8) and an average 96° angle for C–Te–C (entries 11 and 12). The chelating angles, between axial and equatorial positions of type O–Te–C (O1–Te1–C10 and O2–Te2–C20, entries 13 and 14, Table 1) presented an average of 98° angle.

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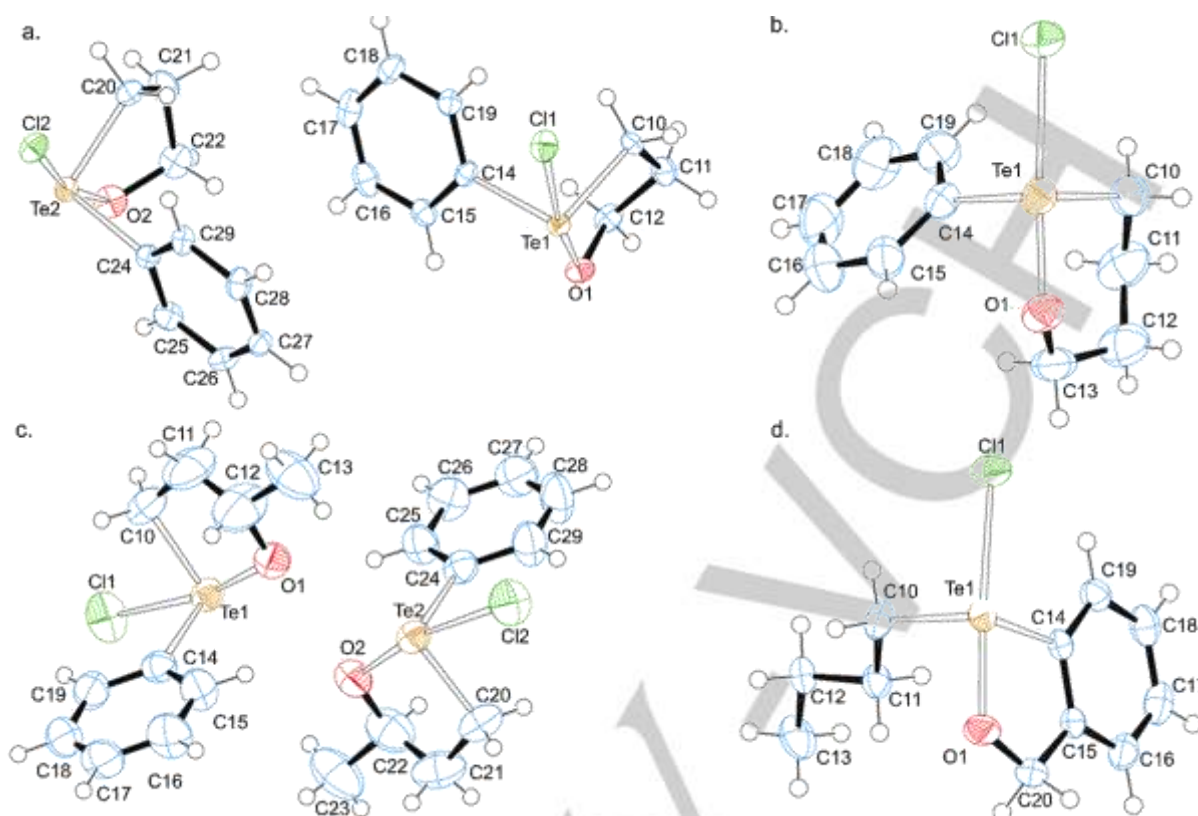


Figure 4. ORTEP^[21] representation of **LQ02** (a), **LQ49** (b), **LQ51** (c) and **LQ56** (d). Thermal ellipsoids are drawn at the 50% probability level. Atom-color: Te-yellow, O-red, Cl-green, C-blue, H-white.

Table 1. Selected Bond Lengths (Å) and Angles (°) for organotelluranes **LQ02**, **LQ49**, **LQ51**, and **LQ56**.

Entry	Compound	LQ02 ^[a]	LQ49	LQ51 ^[a]	LQ56
Bond Lengths (Å)					
1	Te1–O1	2.011(4)	1.983(2)	1.995(4)	2.009(3)
2	Te2–O2	2.000(5)	-	1.993(4)	-
3	Te1–Cl1	2.642(2)	2.6752(9)	2.653(2)	2.6353(11)
4	Te2–Cl2	2.6811(14)	-	2.664(2)	-
5	Te1–C10	2.136(5)	2.134(3)	2.129(5)	2.140(4)
6	Te2–C20	2.144(7)	-	2.118(5)	-
7	Te1–C14	2.132(7)	2.126(3)	2.134(5)	2.111(4)
8	Te2–C24	2.130(7)	-	2.120(6)	-
Angles (°)					
9	O1–Te1–Cl1	170.16(14)	175.68(8)	168.55(11)	170.70(10)
10	O2–Te2–Cl2	165.5(2)	-	169.02(11)	-
11	C10–Te1–C14	94.8(3)	96.12(13)	96.2(2)	98.2(2)
12	C20–Te2–C24	95.5(3)	-	95.6(2)	-
13	O1–Te1–C10	84.5(2)	91.91(13)	83.6(2)	91.5(2)
14	O2–Te2–C20	83.5(2)	-	84.2(2)	-

[a] There are two independent molecules (enantiomers) in the crystal.

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Table 2. Crystal and structure refinement data for organotellurane **LQ02**, **LQ49**, **LQ51**, and **LQ56**.

	LQ02	LQ49	LQ51	LQ56
Empirical formula	C ₉ H ₁₁ ClOTe	C ₁₀ H ₁₃ ClOTe	C ₁₀ H ₁₃ ClOTe	C ₁₁ H ₁₅ ClOTe
Molecular weight (g mol ⁻¹)	298.23	312.25	312.25	326.28
Temperature (K)	173(2)	301(2)	302(2)	200(2)
Crystal system	Orthorhombic	Orthorhombic	Orthorhombic	Monoclinic
Space group	Pca2 ₁	Pbca	Pna2 ₁	P2 ₁ /c
a (Å)	8.6680(4)	10.4460(4)	13.3710(6)	8.4594(6)
b (Å)	9.7474(5)	9.0554(4)	16.9537(9)	16.8778(12)
c (Å)	23.1440(11)	23.4543(11)	10.0889(6)	8.7897(6)
β (°)	90	90	90	108.582(2)
V (Å ³)	1955.45(16)	2218.61(17)	2287.0(2)	1189.54(15)
Z	8	8	8	4
μ (mm ⁻¹)	26.126	2.882	2.796	21.535
θ _{max}	74.979	27.498	27.500	74.957
Reflections collected	136558	170329	150955	70700
Independent reflections	4020	2543	5227	2447
R _{int}	0.084	0.051	0.046	0.076
Reflections with I > 2σ _I	3909	2245	4742	2387
R ₁ (I > 2σ _I)*	0.026	0.027	0.021	0.033
wR ₂ (I > 2σ _I)*	0.066	0.063	0.048	0.093
R ₁ all data ^[a]	0.026	0.033	0.027	0.034
wR ₂ all data ^[a]	0.067	0.065	0.050	0.093
Goodness-of-fit on F ²	1.057	1.154	1.066	1.087
Largest diff. peak and hole (e Å ⁻³)	1.28, -0.60	0.92, -0.81	0.52, -0.40	2.05, -0.81

[a] As defined by the SHELXL program.^[22]

Mechanistic Studies

To further investigate the oxidation-cyclization step for the tellurium-containing heterocycles formation (Scheme 1, step iii), a mechanistic study employing DFT calculations was performed for **LQ02** (see Experimental Section in Supporting Information for computational details). First, the transition state structure (TSS) involved in the oxidation step (**TSS1**) was located, consistent with the chlorine atom transfer from NCS to the Te atom of hydroxy telluride **3a** forming a new Te–Cl bond with a low energy barrier (4.9 kcal mol⁻¹), as shown in Figure 5.

Intrinsic reaction coordinate (IRC) calculations indicated that the Te–Cl bond is 69% formed at the **TSS1** as observed in Figure 6. Moreover, results suggest significant changing of the Cl–Te⋯O angle along this step — ranging from 137° at the reactants to 165° when intermediate is formed (Figure 6) — which was attributed to the formation of strong Te⋯O interaction. Natural Bond Orbitals (NBO) calculations confirmed this hypothesis, revealing electron density donation from oxygen atom to tellurium atom resulting in

a 28.7 kcal mol⁻¹ n(O) → σ*(Te–Cl) interaction, indicating a chalcogen bonding of 2.39 Å^[23]. Results also suggest a highly structural resemblance between protonated intermediate (**I**) and the final product **LQ02**. In fact, the O–H bond is responsible for preventing the formation of **LQ02**, as revealed by comparing Te⋯O and Te–Cl distances and bond lengths: Te–Cl bond increases 0.26 Å from **I** to **LQ02**, whereas the Te⋯O distance was shortened by 0.37 Å after deprotonation, which suggests the formation of a new O–Te–Cl hypervalent bond.

Then we turned our attention to the cyclization step. After several attempts, TSS (**TSS2**) was located. IRC calculations suggest a complex intermediate (**I'**) in the reverse direction, formed by coordination of oxygen atom of succinimide ion (NS⁻) to tellurium atom of **I** (Figure 5 and 6). Since complex **I** is not conformationally appropriate for the formation of **I'**, it is likely that that another NS⁻ anion could be involved in the formation of the latter (coming from reaction occurring nearby), not necessarily the leaving group formed in the oxidation step. After the release of NS⁻, the

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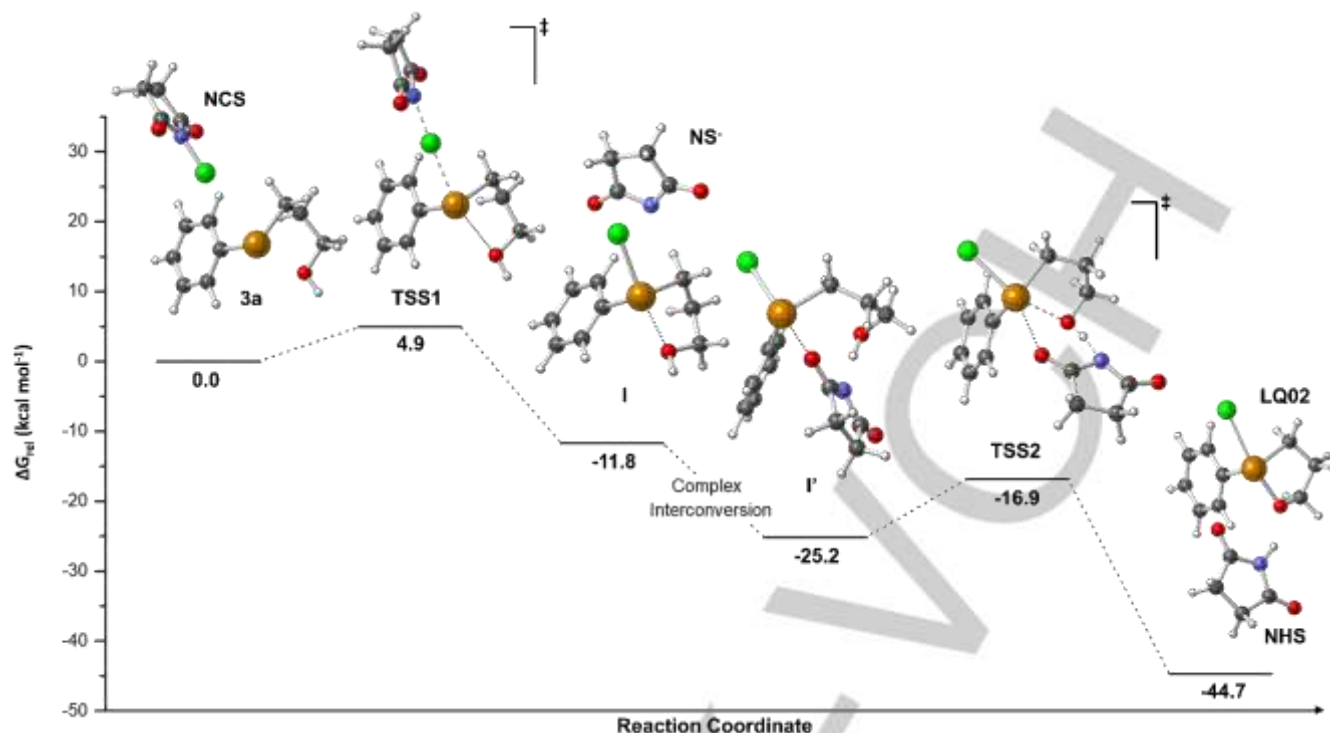


Figure 5. Potential energy surface (PES) of oxidation-cyclization step of the tellurium-containing heterocycles formation obtained using M062X functional combined with LANL2DZ (for Te atom) and 6-311++G(d,p) (for additional atoms). Atom-color: Te-yellow, O-red, Cl-green, N-blue, C-gray, H-white.

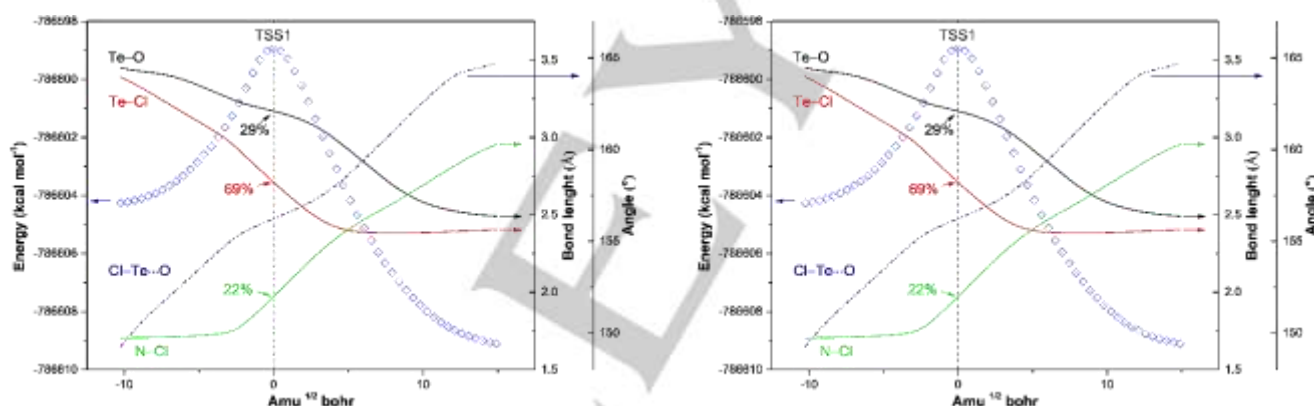


Figure 6. Optimized TSS1 and TSS2 IRC plot obtained with M062X combined with LANL2DZ (Te) and 6-311++G(d,p) basis sets.

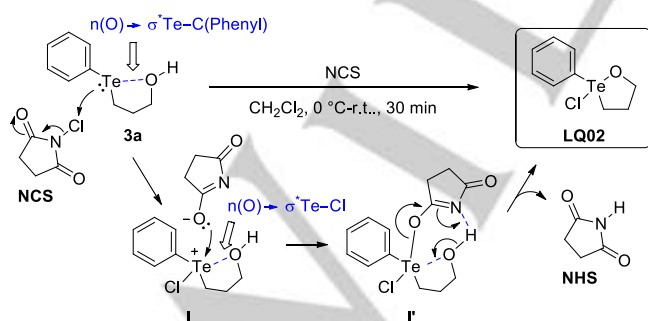


Figure 7. Proposed mechanism for the formation of LQ02.

formation of **I'** is likely to occur in a barrierless process, since a chalcogen interaction ($\text{HO} \cdots \text{Te}$) in **I** is replaced by a $\text{Te}-\text{O}$ covalent bond and an $\text{OH} \cdots \text{N}$ hydrogen bonding or, in other words, a very energetically favorable formation of **I'** should be expected. Moreover, complex **I** features a positive tellurium center, which

makes it a great electrophile. Energy barrier of $8.3 \text{ kcal mol}^{-1}$ to reach **TSS2** from **I'** was observed. Calculations revealed a concerted cyclization step involving the formation of the $\text{Te}-\text{O}$ bond in **LQ02** and the proton transfer from hydroxyl group to the nitrogen atom of NS^- , with elimination of succinimide (**NHS**). The complete mechanistic proposal is presented in Figure 7. IRC plot presented in Figure 6 indicates that 48% of the $\text{Te}-\text{O}$ bond was formed at the **TSS2** whereas 25% of the $\text{Te}-\text{O} \cdots \text{NS}^-$ interaction was broken, consistent with a highly asynchronous pathway. This is also supported by the proton transfer of the $\text{O}-\text{H}$ group to the NS^- anion, which is only 2% completed at the **TSS2**. Uniform $\text{Te}-\text{Cl}$ bond elongation along reaction pathway was also observed. The formation of the $\text{N}-\text{H}$ bond is the last critical event of the cyclization process, finally releasing the products. The oxidation process was found as the rate-determining step and the low barrier of $4.9 \text{ kcal mol}^{-1}$ is consistent with the reaction occurring at 0°C .

Explaining the diastereoselectivity in LQ51 formation

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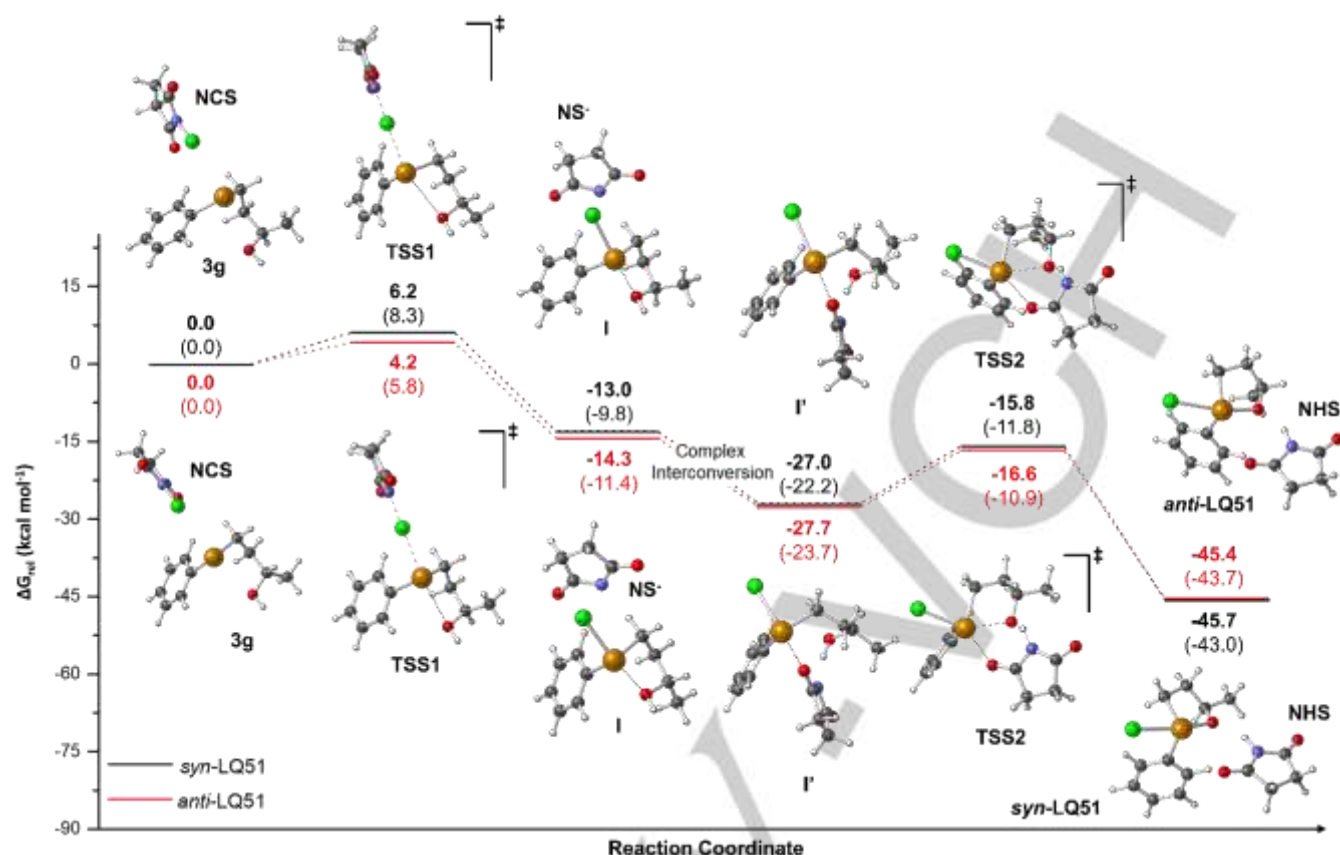


Figure 8. PES of oxidation-cyclization step of the *anti*-LQ51 and *syn*-LQ51 formation obtained using M062X and single points using ω B97X-D (in parentheses) functional combined with LANL2DZ (for Te atom) and 6-311++G(d,p) (for additional atoms). Atom-color: Te-yellow, O-red, Cl-green, N-blue, C-gray, H-white.

1,2 λ^4 -oxatellurane **LQ51** is a two-chirality center-containing compound, so a pair of diastereoisomers is also expected in its preparation. Indeed, the two signals were observed in ¹²⁵Te NMR spectra of **LQ51** at 1101 and 1105 ppm (Figure 3) indicating a highly degree of diastereoselectivity. A 1:4 ratio between diastereoisomers was determined using the relative areas of some equivalent signs at 2.34 and 2.39, 3.50 and 3.61 and 8.16 and 8.21 ppm in the ¹H NMR spectrum (600.13 MHz, CDCl₃, TMS) (see on Figures S88-91 in Supporting Information). Supported by this experimentally observed selectivity and the failure to assign NMR signals to diastereoisomer groups, a mechanistic study employing DFT calculations were also carried out to rationalize the diastereoselectivity and identify the diastereoisomers preferably formed (*syn*-LQ51 or *anti*-LQ51, Figure 3).

TSSs for the oxidation (TSS1) and the cyclization steps (TSS2) were located for each diastereoisomer (*syn*-LQ51 or *anti*-LQ51) as shown in Figure 8. Similar Potential energy surface (PES) and TSSs were obtained for both diastereoisomers of **LQ51** compared to **LQ02** (vide supra). For the oxidation step involved in the formation of *anti*-LQ51, calculations revealed an energy barrier of 4.2 kcal mol⁻¹, which is 2.0 kcal mol⁻¹ lower than that observed for the diastereoisomer *syn*-LQ51 (Figure 8). Such significant difference in the rate determinant step of the reactions corroborates the preferential formation of one of the diastereoisomers, as indicated by NMR analyses. Single point energy calculations using ω B97X-D^[24] were also performed for the stationary points involved in the rate determinant step and similar results were observed. Thus, based on theoretical calculations, *anti*-LQ51 corresponds to the major diastereoisomer formed and observed experimentally in the ¹²⁵Te NMR spectra at

1101 ppm, while the *syn*-LQ51 is the minor diastereoisomer with signal at 1105 ppm.

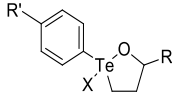
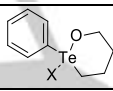
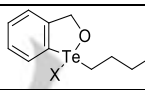
Antiprotozoal Activity of Tellurium (IV)-containing Heterocycles

Hypervalent Te-containing **LQ02-04**, **LQ33-35**, and **LQ48-57** were submitted to *in vitro* assays against *Leishmania amazonensis* promastigotes, and miltefosine was used as the positive control (Table 3). Antileishmanial activities were expressed as IC₅₀ values - the inhibitory concentration able to cause the death of 50% of the parasites in relation to the negative control (non-treated infected cells) - and the cytotoxic of the compounds were expressed as CC₅₀ - the concentration able to kill 50% of the cells (J774A1 macrophages) in relation to the untreated cell control. The selectivity indexes (SI) were determined as the ratio between the CC₅₀ in macrophages and the corresponding IC₅₀ value in promastigotes, allowing to determine how many times the assayed compound was more selective against the protozoa than to the mammalian host cell (macrophages).

1,4-Substituted 1,2 λ^4 -oxatelluranes showed higher activity and selective (expressed by SI) in comparison to unsubstituted analogs **LQ02** (SI: 1, entry 1, Table 2) and **LQ03** (inactive; entry 2, Table 2). In particular, R' = OMe (**LQ04**, SI: 7, entry 3, Table 2) and R' = Cl (**LQ33** and **LQ34**, SI: 5, entries 4 and 5, Table 2) presented higher SI than miltefosine (SI: 3, entry 17, Table 2). **LQ35** (R' = F) exhibited the same value of SI as miltefosine (Figure 9). **LQ48** and **LQ51-55** were considered non-promising compounds due to low SI values observed (≤ 2 ; entries 7-12, Table 2).

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Table 3. *In vitro* Activity toward *L. amazonensis* promastigotes, Cytotoxicity in macrophages, and Selectivity Indexes of **LQ02-04**, **LQ33-35** and **LQ48-57**.^[a]

Entry	Subst.	X	R	R'	IC ₅₀ ± SD (μmol L ⁻¹) ^[b]	CC ₅₀ ± SD (μmol L ⁻¹) ^[c]	SI ^[d]
R'  1,2λ ⁴ -oxatellurolanes:							
1	LQ02	Cl	H	H	34.7 ± 10.6	23.1 ± 2.8	1
2	LQ03	Br	H	H	>100 ^[e]	7.8 ± 2.1	.. ^[f]
3	LQ04	Cl	H	OMe	7.0 ± 1.3	51.3 ± 12.4	7
4	LQ33	Cl	H	Cl	13.0 ± 4.2	64.0 ± 8.3	5
5	LQ34	Br	H	Cl	9.7 ± 0.3	43.7 ± 2.5	5
6	LQ35	Cl	H	F	7.7 ± 2.3	23.3 ± 2.5	3
7	LQ48	Cl	H	Ketal ^[g]	31.4 ± 7.4	20.7 ± 7.4	1
8	LQ51	Cl	Me	H	35.0 ± 3.1	34.0 ± 4.9	1
9	LQ52	Br	Me	H	>100 ^[e]	19.1 ± 2.1	.. ^[f]
10	LQ53	Cl	Ph	H	6.3 ± 0.3	13.9 ± 2.5	2
11	LQ54	Cl	H	Ac	32.8 ± 6.4	13.7 ± 3.7	0.4
12	LQ55	Cl	H	Oxime ^[h]	41.5 ± 7.0	58.0 ± 14.2	1
 1,2λ ⁴ -oxatelluranes:							
13	LQ49	Cl	-	-	8.6 ± 2.5	31.0 ± 7.8	4
14	LQ50	Br	-	-	4.1 ± 1.0	49.0 ± 5.0	12
 1,2λ ⁴ -benzoxatellurolanes:							
15	LQ56	Cl	-	-	5.7 ± 0.3	32.0 ± 6.2	6
16	LQ57	Br	-	-	>100 ^[e]	16.0 ± 2.5	.. ^[f]
17	Miltefosine	-	-	-	20.7 ± 0.2	55.0 ± 1.9	3

[a] Data were expressed as the mean ± SD determined from three independent experiments. [b] IC₅₀: the concentration that inhibit 50% of parasite growth in relation to the untreated control. [c] CC₅₀: cytotoxic concentration able to kill 50% of the macrophages in relation to the untreated control. [d] Selectivity index: CC₅₀/IC₅₀. [e] Considered inactive. [f] Not applicable. [g] 2-Methyl-1,3-dioxolan-2-yl group. [h] (E)-1-(hydroxyimino)ethyl group.

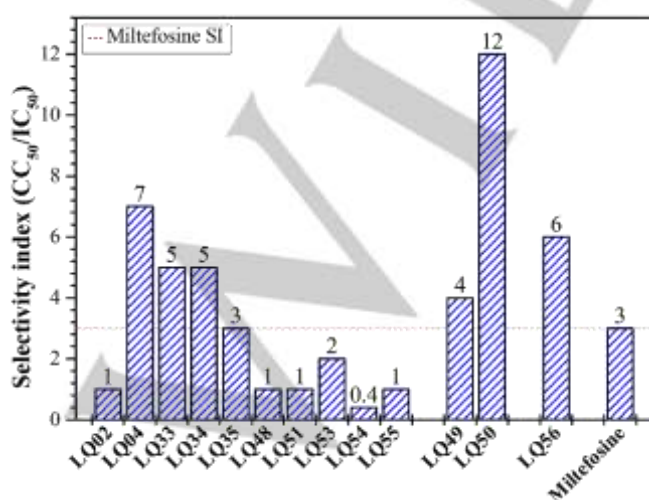


Figure 9. Selectivity indexes of assayed organotelluranes and miltefosine.

The unique two 1,2λ⁴-oxatelluranes assayed here, **LQ49** and **LQ50**, exhibited higher SI than miltefosine (Figure 9). The SI for **LQ49** was 4 (entry 13, Table 2), while **LQ50** was identified as the most selective substance in our series with SI = 12 (entry 14, Table 2). Among 1,2λ⁴-benzoxatellurolanes, only the chlorinated **LQ56** was active (SI: 6, entry 15, Table 2) while its brominated congener **LQ57** was considered inactive, following the trend observed for 1,2λ⁴-oxatellurolanes **LQ02** and **LQ03**. **LQ50** presented the highest SI in the antileishmanial assays (SI = 12, entry 14, Table 2) and evidenced the great potential for Te-containing six-membered heterocycles. The best results observed for 1,2λ⁴-oxatellurolanes and 1,2λ⁴-benzoxatellurolane were highlights in **LQ04** and **LQ56**, respectively. Besides, chlorinated 1,2λ⁴-oxatellurolanes and 1,2λ⁴-benzoxatellurolane showed better performance than equivalent brominated Te-containing heterocycles, except for 1,2λ⁴-oxatelluranes **LQ49** and **LQ50**. Despite the absence of a clear pattern for structure-activity

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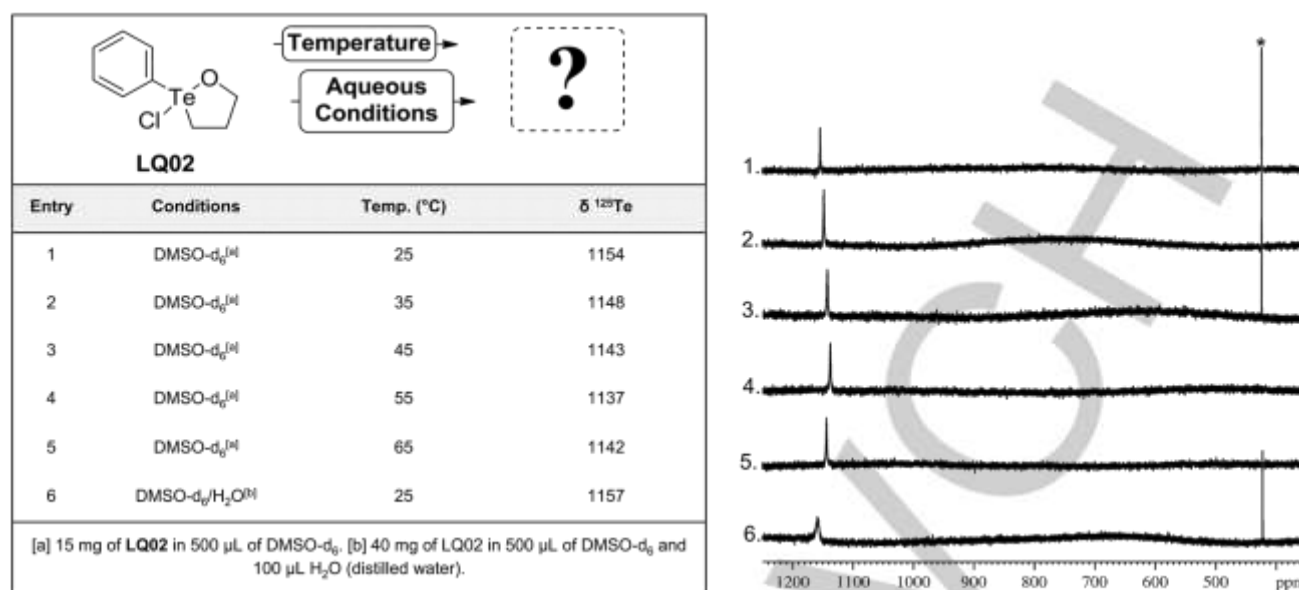


Figure 10. ^{125}Te NMR (126.24 MHz, DMSO- d_6 , PhTeTePh (δ = 422)) on the stability assays for LQ02.

relationships, the three classes of hypervalent Te-containing heterocycles studied in this work - 1,2 λ^4 -oxatelluranes (LQ04), 1,2 λ^4 -oxatelluranes (LQ50) and 1,2 λ^4 -benzoxatellurane (LQ56) - showed potential as antileishmanial agents.

Investigation of Hypervalent Tellurium-containing Heterocycles Stability.

In the literature, there is a debate about the stability of hypervalent tellurium compounds in organic-aqueous media.^[9h, 25] So, complement stability assays for some representative compounds were also performed using ^{125}Te NMR as analytical tool. Initially, the thermal stability of LQ02 in dimethyl sulfoxide- d_6 (DMSO- d_6) was investigated at temperatures ranging from 25 to 65 °C.^[9h, 25a] The results obtained by ^{125}Te NMR are described in Figure 10, in which no significant changes in chemical shifts were observed, demonstrating the thermal stability of the studied organotellurane. Additionally, ^1H NMR analysis confirmed that the structure of LQ02 remained intact. The same methodology was applied for LQ51 to evaluate the possible cycle opening through variations between the proportion of diastereoisomers. However, satisfactorily no structural change was observed even after 5 hours in reflux in CDCl_3 (see on Supporting Information, figure S110).

In the sequence, LQ02 was solubilized in DMSO- d_6 , followed by distilled water addition (Figure 10). ^{125}Te NMR analyses presented a single signal at δ = 1157, again suggesting no chemical structure variation. Recovery and ^1H NMR analysis of LQ02 confirmed no structural modifications in organic-aqueous media. Since the tested organotelluranes did not suffer opening cycle and parallel undesired reactions up to 65 °C, including in organic-aqueous media, as observed for other telluranes,^[24a, b, d] in association with the promising results of biological activity, 1,2 λ^4 -oxatellurane, 1,2 λ^4 -benzoxatellurane and especially 1,2 λ^4 -oxatellurane studied in this work present great potential in the search for biologically active Te-containing compounds.

Conclusion

In summary, eleven chlorinated and five brominated compounds – thirteen unpublished – hypervalent Te-containing heterocycles (LQ02-04, LQ33-35, and LQ48-57) have been successfully synthesized in 0.4% to 40% overall yields and LQ02, LQ49, LQ51, and LQ56 structures were elucidated by NMR analysis and single-crystal X-ray diffraction methods. Computational results revealed that the oxidation-cyclization proceeds via two-step mechanism: (1) the oxidation of the hydroxy telluride by NCS as the rate determinant step with energy barrier of 4.9 kcal mol $^{-1}$ and (2) cyclization involving the formation of the new Te–O bond and the O–H bond break in a concerted asynchronous fashion. Additionally, the diastereoselectivity observed was rationalized using theoretical calculations, allowing the identification of *anti*-LQ51 as the major diastereoisomer formed.

In vitro activity assays against *L. amazonensis* promastigotes showed that 1,2 λ^4 -oxatellurane LQ50 (SI = 12), 1,2 λ^4 -oxatellurane LQ04 (SI = 7) and 1,2 λ^4 -benzoxatellurane LQ56 (SI = 6) were the most promising substances, with SI values higher than that observed for miltefosine. Six-membered 1,2 λ^4 -oxatellurane presented better results than five-membered Te-containing heterocycles (1,2 λ^4 -oxatellurane and 1,2 λ^4 -benzoxatellurane), suggesting this as a more promising class for future investigations. ^{125}Te and ^1H NMR analysis showed LQ02 and LQ51 as thermal stable (up to 65 °C) and in aqueous-organic conditions. In conclusion, this work aims to contribute to the search for bioactive organotellurium compounds and highlight the biological potential of the hypervalent Te-containing heterocycles.

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Keywords: Hypervalent Tellurium compounds • Oxidation-

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Cyclization Mechanism • Leishmanicidal activity • ¹²⁵Te NMR

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Promissory hypervalent Te(IV)-containing heterocycles were obtained with highlights for biological applications. Synthesis, most favorable mechanism identification, *in vitro* antileishmanial potential against *L. amazonensis* and stability assays, supported by ^{125}Te NMR of a series of five- (1,2 λ^4 -oxatellurolanes and 1,2 λ^4 -benzoxatellurols) and six- (1,2 λ^4 -oxatelluranes) membered ring organotelluranes, confirm products containing a Te–O bond.