Synthesis of spiro quasi[1]catenanes and quasi[1]rotaxanes via a templated backfolding strategy

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Due to their well-defined three-dimensional geometry, spiro compounds are widely utilized in drug research. From the central tetrahedral carbon atom, besides the regular structure, an inverted spiro connectivity may be envisioned. Here we disclose the synthesis of this molecule class that we have coined quasi[1]catenanes. Next to their fascinating and aesthetic shape, the higher compactness as compared to regular spiro bicycles is noteworthy. To enable synthetic access to compact entangled multimacrocyclic molecules, we have developed a new strategy. The key element is a template, which is covalently connected to the linear precursors, and spatially directs the sterically congested backfolding macrocyclizations that are required to give quasi[1]catenanes. Similarly, quasi[1]rotaxanes are made.
ntangled molecular architectures inspire synthetic chemists for creative endeavours and find application in nanotechnology and materials research. Striking natural examples are the lasso peptides, and also catenane structures found in some proteins and DNA. Synthetically, an impressive array of entangled molecules such as catenanes and rotaxanes have been made that are mainly applied in nanotechnology research. Supramolecular three-dimensional templating via metal coordination, π-stacking or hydrogen bonding as developed by the respective Sauvage, Stoddart, and Hunter/Vögtle groups gave access to the vast majority of the catenane and rotaxane series. However, the prerequisite of moieties able to form non-covalent interactions limits the structural diversity of the entangled molecules that can be obtained. Remarkably, the first example of a non-statistical catenane synthesis was reported back in 1964 by Schill and Lüttringhaus using a covalent approach. Over the past years, there has been a comeback of covalent approaches towards entangled structures. Currently, we are developing covalent template-directed synthetic concepts to disclose unknown or 'impossible' entangled or mechanically interlocked molecular geometries with access to the natural lasso peptide series as the ultimate goal. Here we report a synthetic concept based on template-directed backfolding macrocyclization of which the utility is demonstrated by the efficient synthesis of an uncommon spiro geometry. The regular spiro geometry is obtained after connection of two rings to the same core tetrahedral carbon atom and can be considered as the molecular equivalent of the figure of eight. The fascinating spirocyclic geometry is found in many natural compounds and due to their rigid and well-defined three-dimensional shape, characterized by a perpendicular arrangement of the two rings, spiro compounds find wide application in drug research. Besides the regular geometry (as for structure 1, Fig. 1), an alternative connectivity may be envisioned by backfolding of the two rings at the central tetrahedral carbon atom to give an inverted spiro configuration (that is, 2, Fig. 1). An intermediate in the landmark catenane synthesis by the Schill group already displayed a similar inverted spiro geometry. Despite the fact that both spiro geometries do not contain any stereocentre, they are still interesting from a stereochemical point of view. Uniquely, in the spiro connectivity case, after inverting the centre-of-symmetry tetrahedral carbon, a diastereomer is obtained with different properties while retaining the $C_2$ symmetry around the same axis. Although we realize that every name is a compromise because no mechanical bond is present, we coined the topologically trivial inverted spiro geometry a quasi[1]catenane (ref. 25). Similarly, a quasi[1]rotaxane 3 can be drawn, of which unwinding to give conformer 4 is prevented by large stoppers (Fig. 1).

For steric reasons, the synthesis of the quasi[1]catenane and quasi[1]rotaxane architectures requires specifically tailored methodologies. We have tackled this challenge, as will be pointed out first for quasi[1]catenane 2, by using a covalent template directing the required backfolding cyclization. Template 6 (Fig. 2, route A) will be temporarily connected to two of the four linear ring precursors in 5 and will eventually be part of the spirocyclic ring formed with the other precursor chains. Due to the tetrahedral geometry of the central carbon atom, cyclization of the first ring on the template in 7 will occur in a perpendicular and backfolded fashion over the linear precursor of the second ring to form a pseudo-quasi[1]rotaxane 8 (ref. 26). Subsequent backfolding cyclization of the second ring to give the inverted spiro connectivity, followed by cleavage of the temporal scaffolding bonds provides bicyclic quasi[1]catenane 2. Essentially, spiro quasi[1]catenane 2 may be obtained from the tetrahedral precursor 5 in just four steps, of which three are macrocyclizations. Without making the temporal connections to the template, the regular spiro bicycle 1 will be obtained (Fig. 2, routes B and C). An alternative route to the regular spiro bicycle 1 opens up after breaking the temporary bonds in 8 (Fig. 2, route D), initiating unwinding of the sterically congested quasi-thread fragment to give 10, followed by closure of the second ring. Capping the end-groups in pseudo-quasi[1]rotaxane 8 by large stoppers (Fig. 2, route E) to give 12 prevents the unwinding process after temporal bond cleavage to give quasi[1]rotaxane 3. Stopper attachment to 10 (Fig. 2, route F) gives the sterically relaxed conformer 4.

**Results**

**Design of the building blocks.** To avoid the requirement of protective groups and to facilitate the sterically difficult macrocyclizations, robust transformations were selected, that is, transesterification/lactonization for the scaffolding temporal connection to the template, Cu-catalysed azide alkyn cyclodaddition (CuAAC) for the first ring closure and olefin metathesis for the second and final ring closure or introduction of the quasi-rotaxane stopper elements. For synthetic reasons and to ensure the perpendicular mutual arrangement of the two pairs of ring-precursor chains at precursor 5, we have chosen a spiro-linkage via the tetrahedral carbon atom of a 9H-fluorene moiety (Fig. 3). The design of the temporal covalent scaffolding tether at the central precursor 5 is based on amide-N-benzylid moieties containing phenolic hydroxyl groups that will be esterified to template 6 and may be removed from the final spiro or rotaxane compounds by consecutive transesterification and protolytic cleavage of the amide benzylid linkages. As the template, 2,5-bis(pent-4-yn-1-yl)oxy)-1,4-benzendicarboxylic ester 6 was chosen. The tether length in precursor 5 and template 6 gives access to spiro architectures of 27- and 31-membered rings, resulting from CuAAC and ring-closing metathesis (RCM).
Synthesis of the quasi[1]catenane. We first followed the strategy to quasi[1]catenane 2 incorporating the inverted spiro[27,31] framework (Fig. 2, route A). Connection of 5 via transesterification of the pentafluorophenol ester 6a using Cs$_2$CO$_3$ as the base in acetonitrile (2 mM) gave the 21-membered macrocyclic bislactone 7 in 69% isolated yield (Fig. 4a). The subsequent backfolding double CuAAC reactions required careful optimization to avoid competing intermolecular reactions over the challenging process to the 31-membered cage-type macrocycle. Slow addition of a 5 mM solution of 7 to a solution of Cul (2 equiv) and DIPEA/lutidine (4 equiv) in acetonitrile gave 8 in poor and irreproducible yields due to extensive insoluble polymer formation. Simply stirring a 5 mM dichloromethane solution of 7 at reflux for 24 h using tetrakis(acetonitrile)copper(I) tetrafluoroborate (0.25 equiv) and tris(1-benzyl-1H-1,2,3-triazol-4-yl)methylamine (TBTA, 0.25 equiv) as the catalytic complex gave, after work-up and purification, 8 in a 79% reproducible yield. The final backfolding closure of the 27-membered cycloalkene via RCM using Grubbs’ second-generation catalyst at high dilution (1 mM) in dichloromethane at reflux gave the tricyclic product in an isolated yield of 71% as a mixture of E/Z isomers. As a coeluting side product, the liquid chromatography mass spectrometry trace also showed some presence of the 26-membered cycloalkene analogue, due to double bond migration before cyclization. After removal of the temporary tether by consecutive transesterification using NaOMe and trifluoroacetic acid (TFA)/Et$_3$SiH-mediated protolytic cleavage of the benzylic linkages, the quasi[1]catenane 1 was isolated in a last-three-steps yield of 66%. $^1$H NMR studies confirmed the solution phase C2-symmetric conformation of the inverted spiro quasi[1]catenane 2 as drawn in Fig. 4a. Most protons of the nine pairs of cycloalkene methylene protons gave well-separated signals. In addition, the strong nOe contacts between the fluorene 8 and 8’ protons and the majority of the aliphatic protons of the macrocycloalkene confirm the folding of the macrocycle over the fluorene moiety (see Supplementary Information). Fortunately, after catalytic hydrogenation to remove the alkene E/Z mixture, crystals of 2-2H$_2$ could be obtained of sufficient quality for X-ray diffraction (Fig. 4b). Besides unequivocal proof of the backfolded quasi[1]catenane configuration, the solid-state structure showed a similar conformation as in solution, albeit with the flexible C20-macrocyloalkene offset folded over the fluorene moiety.

Next, the regular spiro[27,31] macrobicycle 1 was made. Ironically, initial attempts were troublesome. Subjection of 5 to the high-dilution RCM macrolactonization conditions (Fig. 2, route B) led to complex and inseparable mixtures. Fortunately, by beginning with the CuAAC reaction from 5 and 6b,
Figure 4 | Synthetic routes from the common quaternary carbon precursor 5. (a) Detailed reaction conditions for the synthesis of spiro bicycle 1 and quasi[1]catenane 2: (i) Cs2CO3 (4 equiv), MeCN (2 mM), 4 Å molecular sieves, 60 °C. (ii) Cu(MeCN)4BF4 (0.25 equiv), TBTA (0.25 equiv), CH2Cl2 (5 mM), reflux. (iii) Grubbs second-generation catalyst (0.2 equiv), CH2Cl2 (1 mM), 30 °C. (iv) NaOMe (5 equiv), MeOH/THF (1:1). (v) TFA/CH2Cl2 (9:1), Et3SiH (20 equiv). (vi) HCl (3 M), MeOH. (b) Molecular structure of quasi[1]catenane 2-H2 in the crystal. Hydrate water and the disorder present in the C20 aliphatic chain have been removed in the drawing. (c) Detailed reaction conditions for the synthesis of quasi[1]rotaxane 3 and its un wound conformer 4: (vii) Grubbs second-generation catalyst (0.2 equiv), CH2Cl2 (5 mM), reflux. (viii) HCl (3 M), MeOH/THF (4:1). Further conditions see a. TBTA = tris[1-(1-benzyl-1'H-1,2,3-triazol-4-yl)methyl]amine, Grubbs second-generation catalyst = (1,3-Bis(2,4,6-trimethylphenyl)-2-imidazolidinylidene) dichloro-(phenylmethylene)(tricyclohexylphosphine)ruthenium.

Monocycle 10 was obtained, although in a yield of 42% only (Fig. 4a, route C). It should be noted that this yield is significantly lower than the CuAAC reactions to 8 that both occurred in a template/scaffold-preorganized intramolecular fashion. As an elegant detour, monocycle 10 could be obtained quantitatively after methoxide-induced cleavage of the tether in pseudo-quasi[1]rotaxane 8 inducing immediate steric relaxation (Fig. 4a, route D). By lowering the temperature to 30 °C at high dilution, the RCM reaction proceeded cleanly and no truncated cycloalkene was observed. Protolytic removal of the 2-hydroxy-4-methoxybenzyl groups led to an unexpectedly stable adduct of TFA and 1, which could not be separated. However, treatment with methanolic 3 M HCl at 50 °C gave a clean removal of the benzyl groups and the resulting triazolium ions were neutralized with aqueous NaHCO3 to give the regular spiro macrocycle 1 in a yield of 51%. Catalytic hydrogenation removed the RCM-derived olefin E/Z mixture to give 1-H2. The different geometries of 1-H2 and 2-H2 are obvious from the 1H NMR spectra (Fig. 5) showing almost no overlapping signals. In quasi[1]catenane 2 especially, the four pairs of methylene protons (Fig. 5, protons f–i) experience an upfield shift to the 0.5 p.p.m. region due to their presence in the shielding region above the plane of the aromatic fluorene phenyl groups. For the same reason, a dramatic upfield shift of over 3 p.p.m. of the NH triplets (Fig. 5, protons k) of 2 is observed. The inverted spiro geometry in 2 is also reflected by the hydrogen bonds between the triazole CH’s (Fig. 5, protons s) and the amide carbonyls resulting in a 0.8 p.p.m. downfield shift as compared to the same protons in regular spiro compound 1. In addition, donor–acceptor distances in the crystal structure of 2.2 and 2.4 Å, respectively, show these hydrogen bonds in the solid...
Figure 5 | ¹H NMR spectra. Overlays of the ¹H NMR spectra clearly show the similarities between the geometries of quasi[1]catenane 2-H₂ and quasi[1]rotaxane 3, and also between spiro bicycle 1-H₂ and monocyke 4. Furthermore, the structural dissimilarity between both C₂-symmetric but diastereomeric spiro bicycle 1 and quasi[1]catenane 2-H₂ is reflected by their spectra as is the case for quasi[1]rotaxane 3 and its unwound conformer 4.

Synthesis of the quasi[1]rotaxane. The backfolded intermediate 8 also provides access to quasi[1]rotaxane 3 (Fig. 4c). Cross-metathesis³⁹ of the terminal thread alkenes in 8 with the acrylamide-functionalized stopper 11, followed by tether detachment by consecutive transterification and protolysis gave quasi[1]rotaxane 3 in a yield of 39% over three steps. The unthreaded conformation 4 was obtained via cross-metathesis-mediated tether attachment to unwind intermediate 10 and protolytic debenzylisation in a two-step yield of 28%. The structural dissimilarity between the quasi[1]rotaxane 3 and conformer 4 was reflected by comparing the ¹H NMR spectra showing, as in the quasi[1]catenane series, remarkable shift differences, especially for the methylene protons in the thread fragment close to the fluorene moiety, the central amide NH’s and the triazole proton (Fig. 5). Quasi[1]rotaxane 3 and the unwound isomer 4 only differ in conformation of which interconversion is mechanically blocked by the stoppers. In contrast to the very facile unwinding of the pseudo-thread in 8 through the 31-membered ring to give 10 (Fig. 4a, route D), the tris(4-(tert-butyl)phenyl)methane stoppered quasi[1]rotaxane 3 conformation was completely stable, even after heating in dimethylsulphoxide at 110°C for 18 h as was confirmed by ¹H NMR spectroscopy.

Discussion
Generally, spiro compounds are obtained after two ring closures of the four linear-chain ring precursors connected to a central tetrahedral carbon atom. We have shown that after prior temporal covalent connection via a template of two of the four ring-precursor chains, ring closure of the second pair to the template is forced in a backfolded fashion. Final closure of the first pair of ring-precursor chains followed by cleavage of the temporary linkers to the template provides an inverted spiro geometry coined here as a quasi[1]catenane. Although the regular spiro and quasi[1]catenane geometries only differ by inversion of the configuration of the centre-of-symmetry tetrahedral carbon, they relate as diastereomers with inherently different physical properties as reflected by their contrasting ¹H NMR spectra. By starting from the same tetrahedral carbon-centred acyclic precursor and the same backfolding concept, the structurally related mechanically locked quasi[1]rotaxane and its unwound conformer were obtained. The next step in the backfolding cyclization concept will be replacing the permanent perpendicular spiro connectivity by structurally similar but cleavable moieties, such as a 1,3-dioxolane, to give 2[rotaxanes and [2]catenanes. The longer-term goal is to apply the covalent tether-directed backfolding cyclization concept to synthetically disclose the natural [1]rotaxane-type lasso peptide series by incorporation of two amides of the thread section of the peptide backbone into a imidazolizin-4-one as the perpendicular and cleavable thread/ring connecting moiety. Covalent linking in a perpendicular fashion of the ring over thread fragment, as is the case with the backfolding cyclization concept, ensures the atom-precise mutual positioning of these components, which is a prerequisite for the eventual total synthesis of the lasso peptides.

Methods
Macrolactonization of the tetrahedral precursor 5 to template 6a. 5 (1.94 g, 1.89 mmol), 6a (1.38 g, 2.08 mmol, 1.1 equiv), Cs₂CO₃ (2.45 g, 7.58 mmol, 4 equiv) and 4 Å molecular sieves (1.00 g) were dissolved in dry CH₃CN (750 ml) and the mixture was stirred overnight at 60°C under a N₂ atmosphere. The solvent was evaporated and the residue was taken in ca. 20 ml CH₂Cl₂ and filtered through a plug of Celite, which was washed with CH₂Cl₂. The organic layer was concentrated in vacuo and dry-loaded on silica and purified by column chromatography (petroleum ether (PE):EtOAc 4:1→3:1) to give 7 (1.72 g, 1.31 mmol, 69%) as a thick colourless oil.

state of 2. The different chromatographic retention times of the two spiro diastereomers visualizes the different physical properties showing a higher polarity for the densely packed quasi[1]-catenane 2.

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Double CuAAC macrocyclizations towards cage 8. 7 (420 mg, 0.319 mmol) and TBT (42 mg, 0.080 mmol, 0.25 equiv) were dissolved in 65 ml dry CH₂Cl₂ and degassed with 5 vacuum/N₂ cycles. After which CuCl(CH₃CN)BF₄ (25 mg, 0.080 mmol, 0.25 equiv) was added and the mixture was stirred overnight at reflux under N₂ atmosphere. The reaction mixture was concentrated in vacuo and dry-loaded on silica and purified by column chromatography (PE:EtOAc 1:1→1.2→0.1) to give 8 (333 mg, 0.253 mmol, 79%) as a colourless foam.

Ring-closing metathesis to the pseudo-[quasi]-1:catenane. 8 (79 mg, 0.060 mmol) was dissolved in dry CH₂Cl₂ (60 ml) and degassed with 5 vacuum/N₂ cycles. To the solution was added Grubs second-generation catalyst (10 mg, 0.012 mmol, 0.2 equiv) and the mixture was stirred overnight at 40°C. The mixture was concentrated in vacuo, dry-loaded on silica and purified by column chromatography (PE:EtOAc 1:2→1.3→1.4) to give the pseudo-[quasi]-1:catenane (55 mg, 0.0427 mmol, 71%) as a beige solid.

Temporary tether removal to liberate quasi-[1]:catenane 2. The pseudo-[quasi]-1 catenane (55 mg, 0.043 mmol) was dissolved in dry THF/CH₂OH (2 ml, 1:1) and anhydrous NaOCH₃ (12 mg, 0.21 mmol, 5 equiv) was added and the mixture was stirred at room temperature for 1 h. The reaction was quenched by addition of 0.5 ml AcOH and the reaction was diluted with 15 ml EtOAc and 10 ml saturated NaHCO₃. The water layer was extracted with 2 × 5 ml EtOAc and the obtained organic layers were dried over MgSO₄ and concentrated in vacuo to give the dimethyl ester (58 mg, 0.0427 mmol, quantitative) as a slight yellow film. To this, TFA/CH₂Cl₂ (5 ml, 9:1) and Et₃SiH (0.136 ml, 0.854 mmol, 20 equiv) were added and the mixture was stirred overnight at room temperature and concentrated in vacuo. The residue was dissolved in CH₂Cl₂ (10 ml) and Et₃N (0.5 ml) was added and stirred for 5 min. The organic layers were washed with 1 M HCl (10 ml), saturated NaHCO₃ (10 ml), dried over MgSO₄ and concentrated in vacuo. The crude product was dry-loaded on silica and purified by column chromatography (CH₂Cl₂/CH₂OH 96:4→94:6) to give 2 (43 mg, 0.0404 mmol, 93%) as a colourless foam.

Data availability. Additional data supporting the findings reported in this article are available within the Supplementary Information file. For the experimental procedures and spectral data of all compounds described, see the Supplementary Methods. For 1H and 13C NMR spectra of all compounds, see Supplementary Information.

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