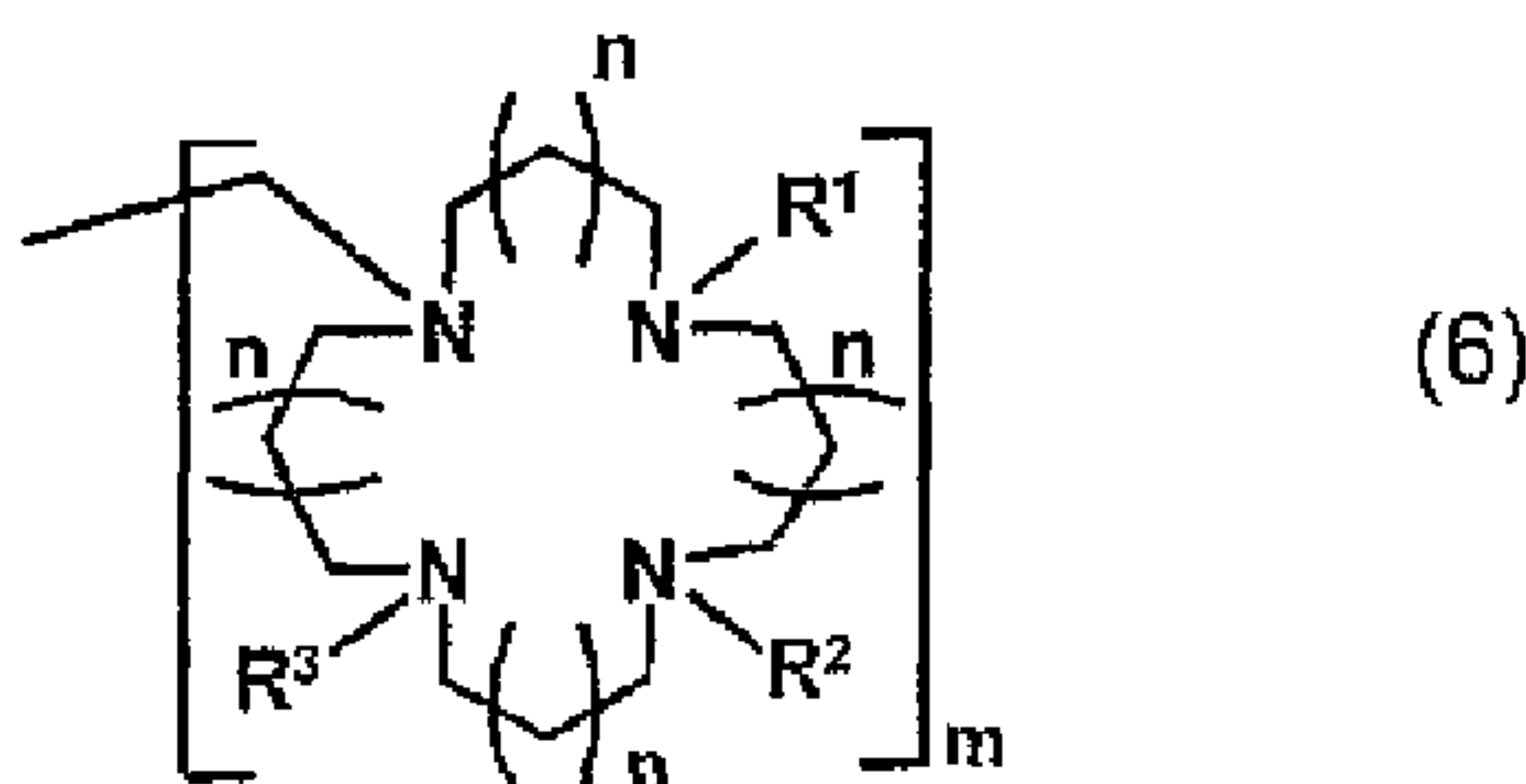
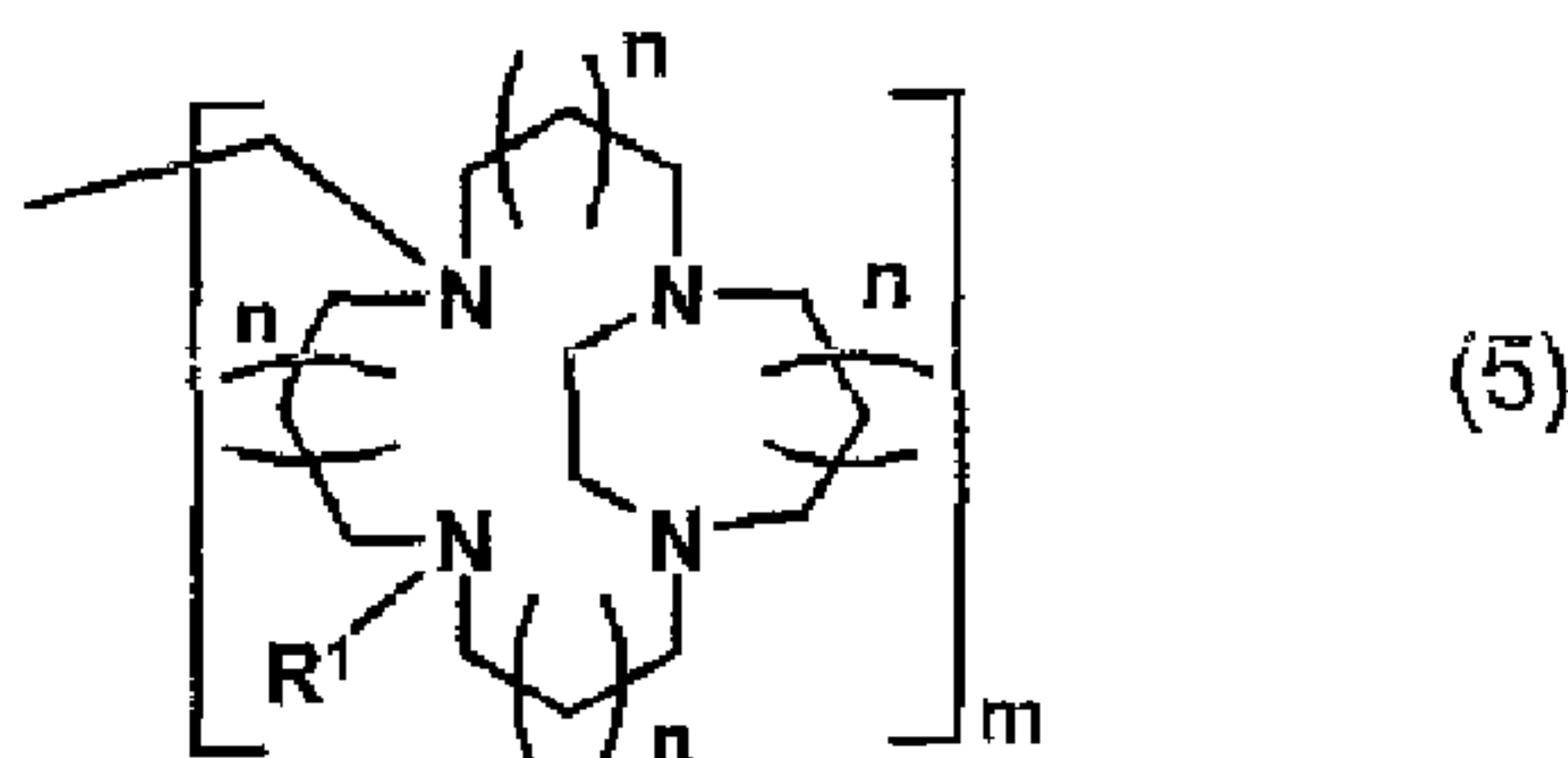
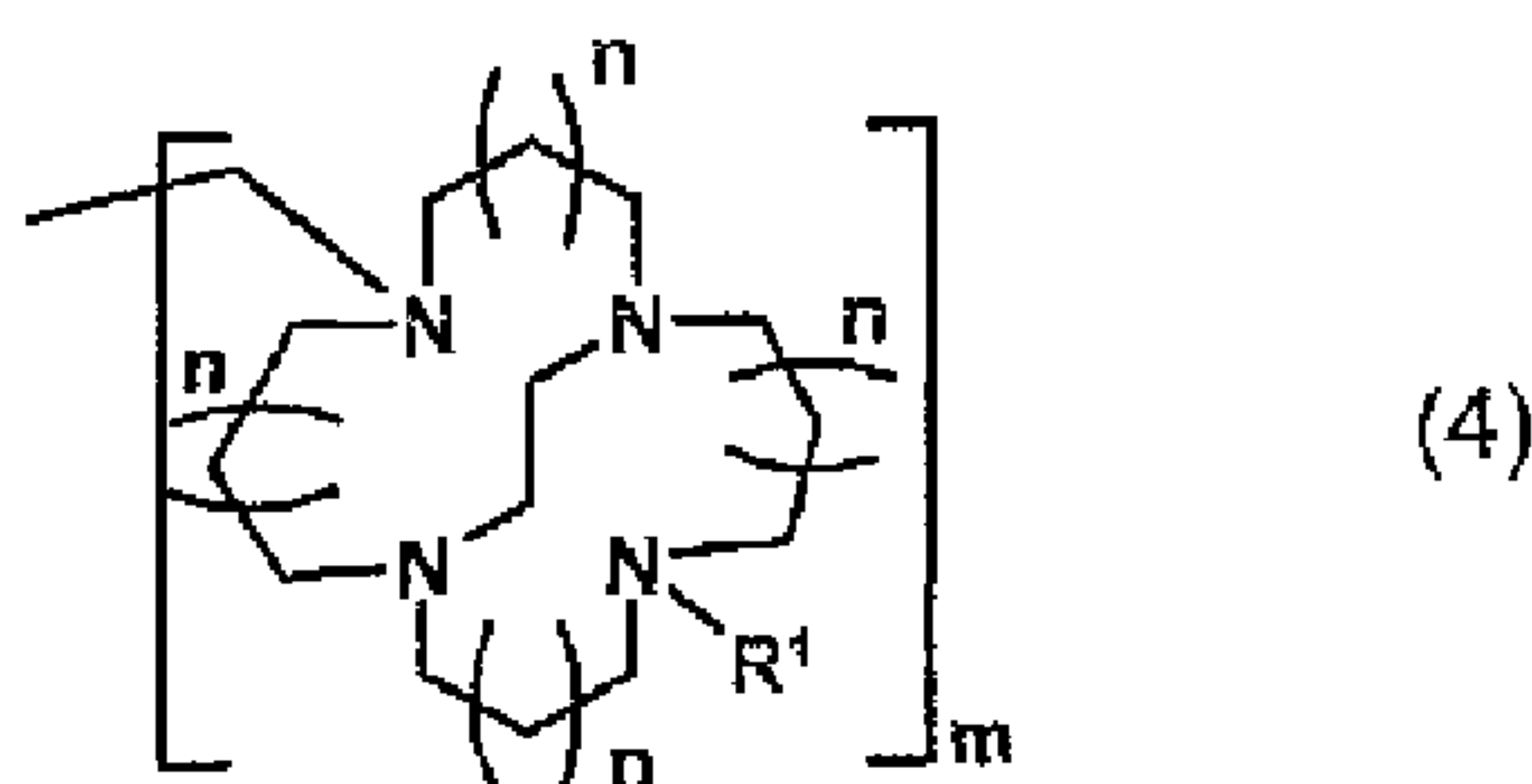




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(54) Titre : NOUVEAUX DERIVES DE MACROCYCLES ET COMPLEXES METALLIQUES ANTIVIRAUX COMPRENANT
DES MACROCYCLES PONTES
(54) Title: NOVEL ANTIVIRAL MACROCYCLE DERIVATIVES AND METAL COMPLEXES, INCORPORATING BRIDGED
MACROCYCLES



(57) Abrégé/Abstract:

A compound of Formulae (1), (2) or (3), A₁-Ph (1) or A₁-Ar-A₂ (2) or A₁-(A₂)A'-A₃ (3) wherein A₁, A₂ and A₃ are selected from Formulae (4), (5) and (6), or a salt whereof; wherein Ar is selected from the group consisting of 1,2-phenyl, 1,3-phenyl and 1,4-



(57) **Abrégé(suite)/Abstract(continued):**

phenyl optionally substituted with one or more methyl groups; m and n are independently integers from 0 to l; Ph is phenyl, optionally substituted with one or more methyl groups; Ar^l is trisubstituted phenyl optionally substituted with one or more phenyl groups; and R¹, R² and R³ are independently selected from the group consisting of H, Me and linear chain C₂- C₄ alkyl.

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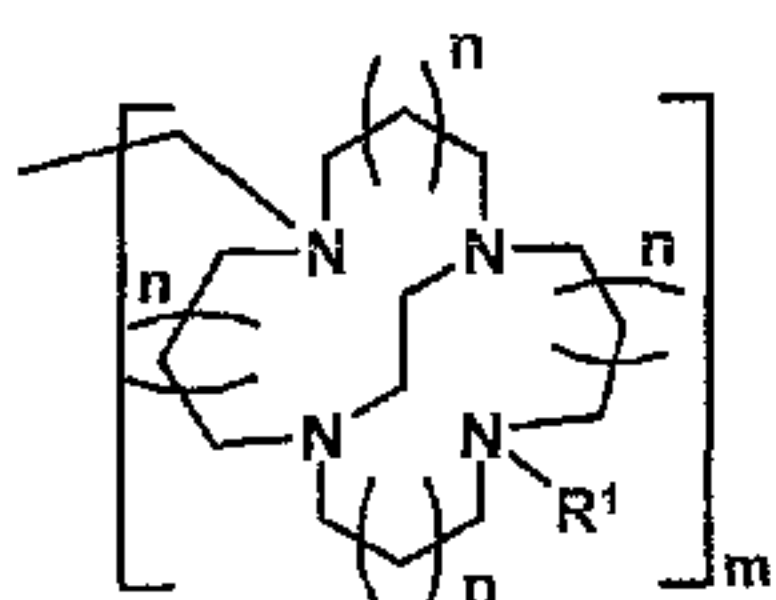
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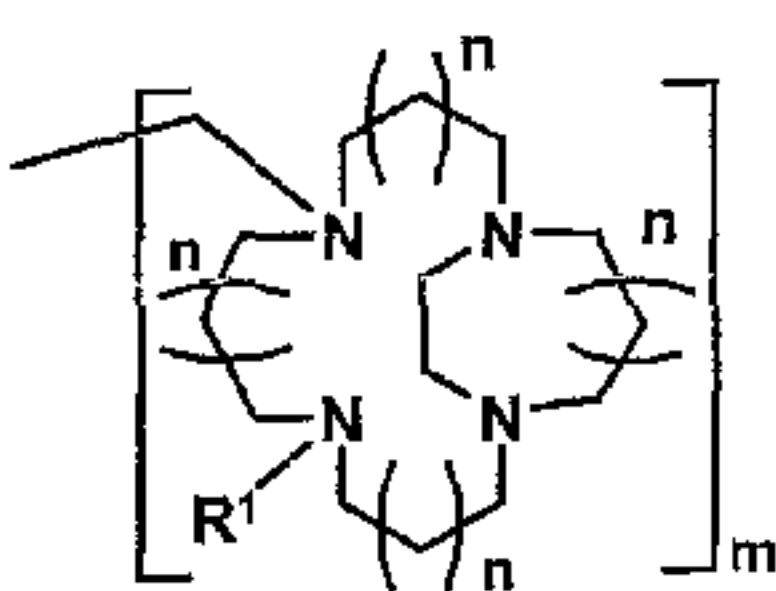
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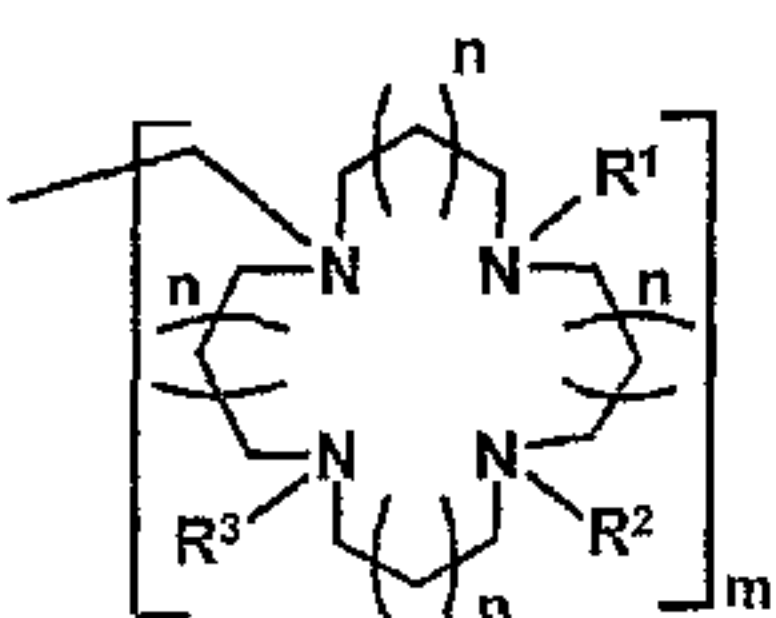
(54) Title: NOVEL ANTIVIRAL MACROCYCLE DERIVATIVES AND METAL COMPLEXES, INCORPORATING BRIDGED MACROCYCLES



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(57) Abstract: A compound of Formulae (1), (2) or (3), A₁-Ph (1) or A₁-Ar-A₂ (2) or A₁-(A₂)A'-A₃ (3) wherein A₁, A₂ and A₃ are selected from Formulae (4), (5) and (6), or a salt whereof; wherein Ar is selected from the group consisting of 1,2-phenyl, 1,3-phenyl and 1,4-phenyl optionally substituted with one or more methyl groups; m and n are independently integers from 0 to 1; Ph is phenyl, optionally substituted with one or more methyl groups; Ar¹ is trisubstituted phenyl optionally substituted with one or more phenyl groups; and R¹, R² and R³ are independently selected from the group consisting of H, Me and linear chain C₂-C₄ alkyl.

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NOVEL ANTIVIRAL MACROCYCLE DERIVATIVES AND METAL COMPLEXES, INCORPORATING BRIDGED MACROCYCLES

FIELD OF THE INVENTION

5 This invention relates to novel mono -or poly-azamacrocycle derivatives incorporating bridged macrocycles, methods of production thereof and dosage forms incorporating these derivatives. The invention also relates to use of the derivatives for manufacture of medicaments and to a method of treatment using these dosage forms. Preferred compounds in accordance with this invention are useful as CXCR4 co-
10 receptor antagonists, especially for anti-HIV and other anti-viral treatments.

BACKGROUND OF THE INVENTION

 CXCR4 chemokine receptors, are found on the surface of immune cells, and interact with the specific natural ligand, known as CXCL12 or stromal cell-derived
15 factor 1 α (SDF-1 α). They have been revealed to play a role in a number of disease states. For example the CXCR4 SDF-1 α system has involvement in cancer progression and metastasis, and in the development of rheumatoid arthritis. CXCR4 and CCR5 co-receptors have been identified as the entry route for HIV into cells, providing a new therapeutic approach to treatment by entry inhibitor drugs rather than
20 the current preference for reverse transcriptase and protease inhibitors. An object of the present invention is to provide new antagonists for the CXCR4 co-receptor.

 This invention may provide novel compounds that selectively bind to chemokine receptors based on cross bridged macrocycles. The compounds may incorporate metal
25 ions and form stable complexes prior to administration.

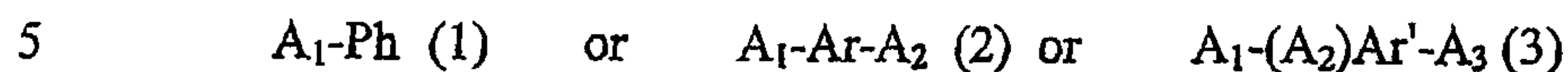
 It is an object of this invention to provide examples of bridged azamacrocyclic derivatives that show specific chemokine receptor binding.

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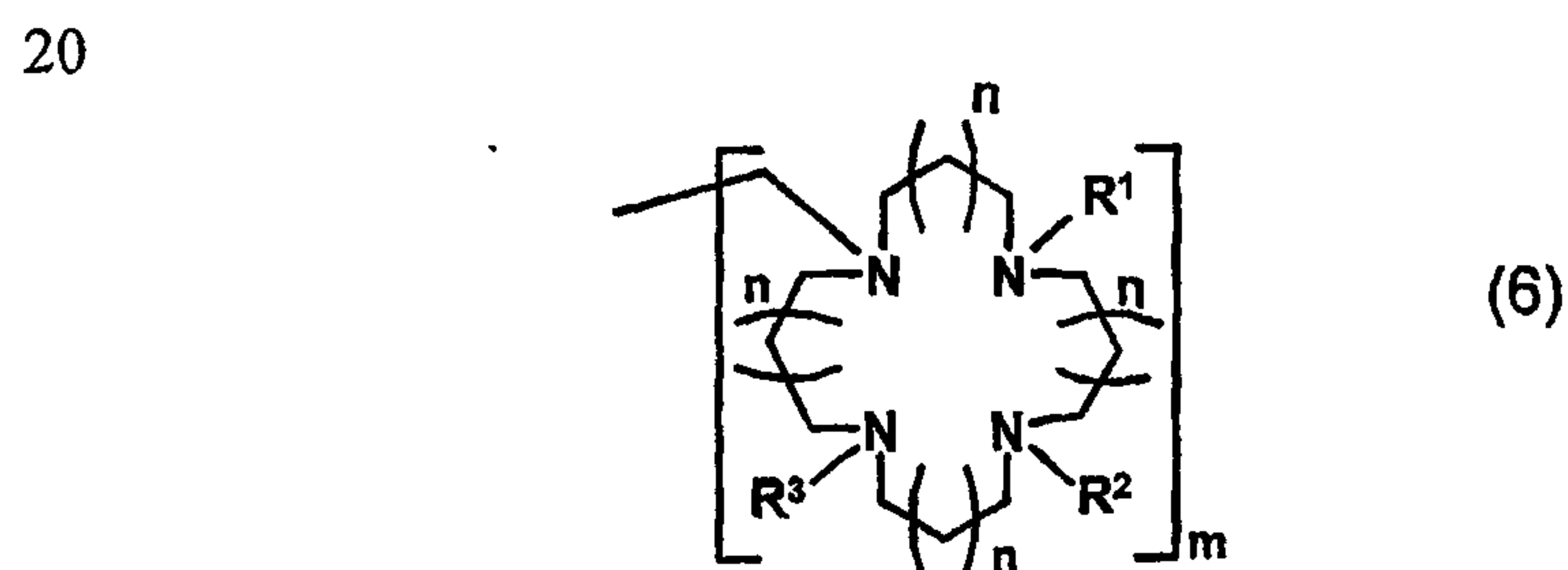
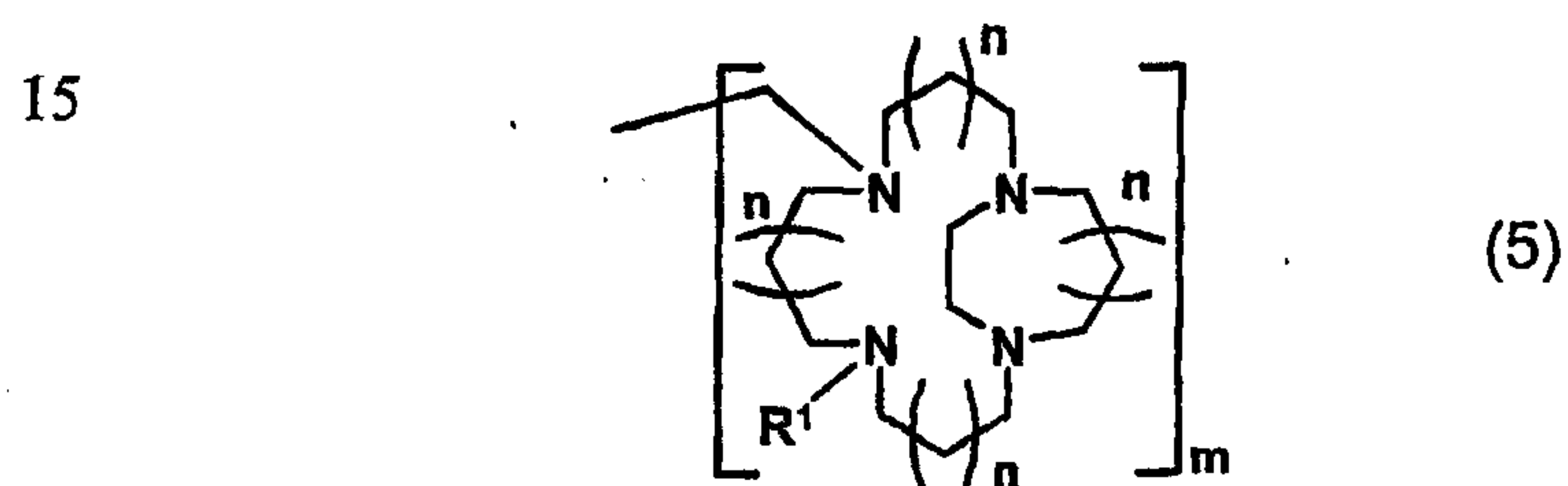
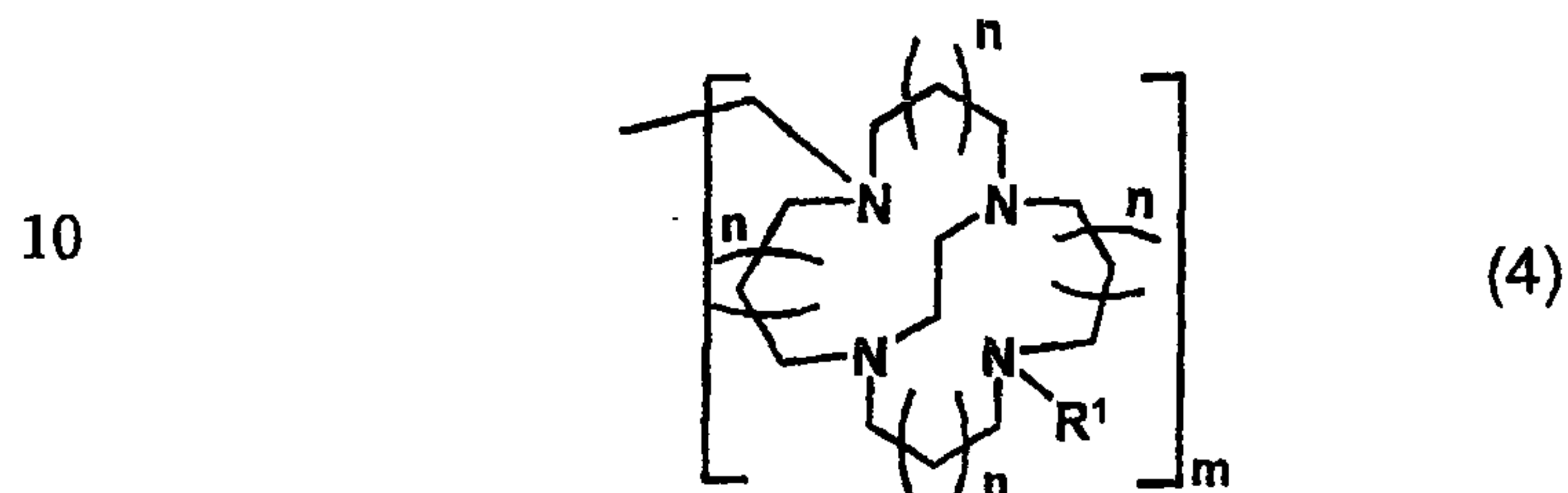
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SUMMARY OF THE INVENTION

According to a first aspect of the present invention there is provided compound of Formulae 1, 2 or 3,



wherein A_1 , A_2 and A_3 are selected from Formulae 4, 5 and 6:



or a salt whereof;

wherein Ar is selected from the group consisting of 1,2-phenyl, 1,3-phenyl and 1,4-phenyl optionally substituted with one or more methyl groups;

30 m and n are independently integers from 0 to 1;

Ph is phenyl optionally functionalised with one or more groups R^4 , wherein R^4 is selected from the group consisting of: methyl, other $C_1\text{-C}_4$ alkyl and fluorophores;

Ar' is trisubstituted phenyl optionally substituted with one or more phenyl groups; and

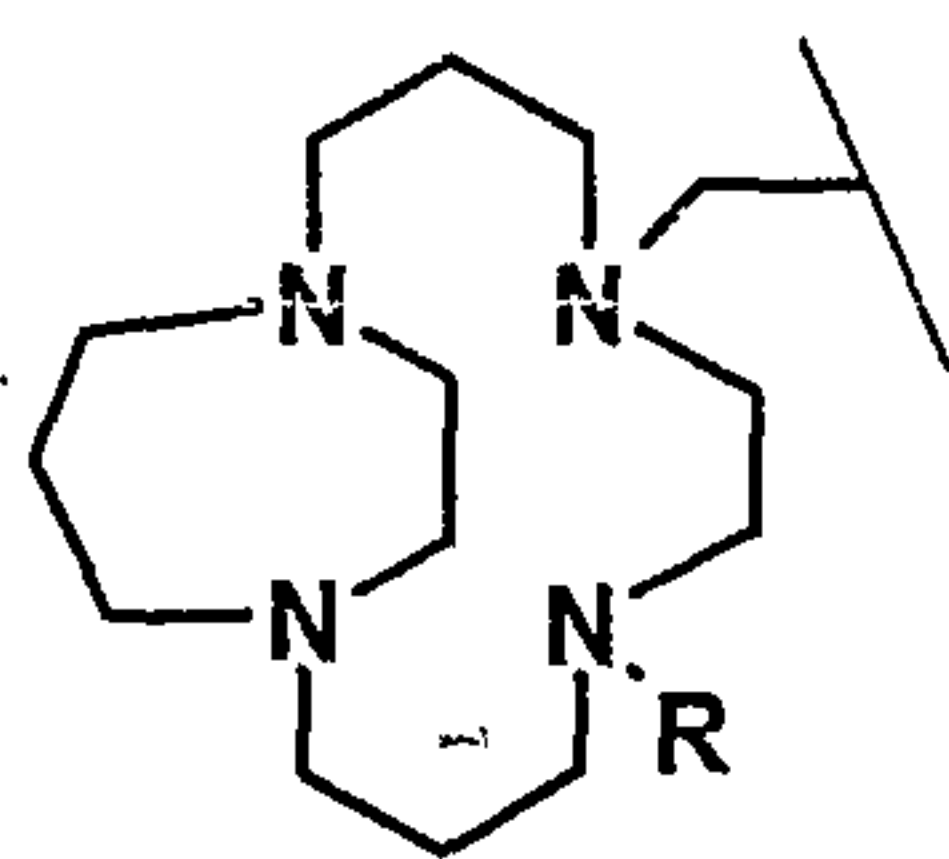
R¹, R² and R³ are independently selected from the group consisting of H, Me and linear chain C₂- C₄ alkyl;

5 with the provisos that when Ph is unsubstituted phenyl in Formula 1 and A₁ is Formula (4), R₁ is not Me; and

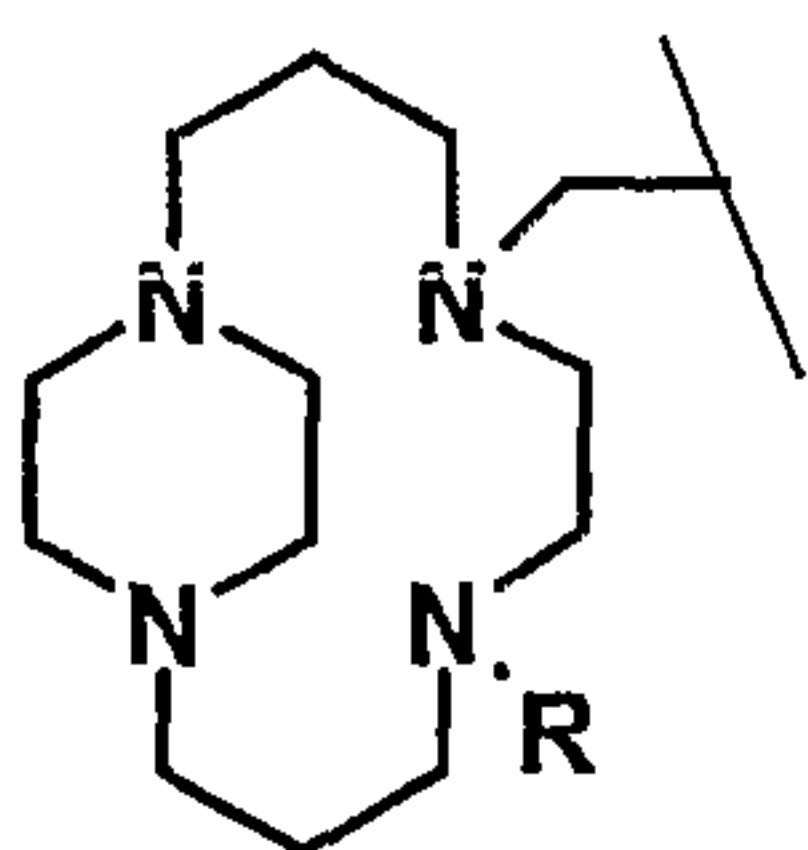
wherein not more than one of A₁ and A₂ in Formula (2) or no more than two of A₁, A₂ and A₃ in Formula (3) is Formula (6).

10 The invention further provides a complex of a compound selected from Formulae 1,2 or 3 with a metal ion selected from the group consisting of: transition elements and mixtures thereof. In preferred compounds the complex is formed with divalent ions of copper and zinc or a mixture thereof.

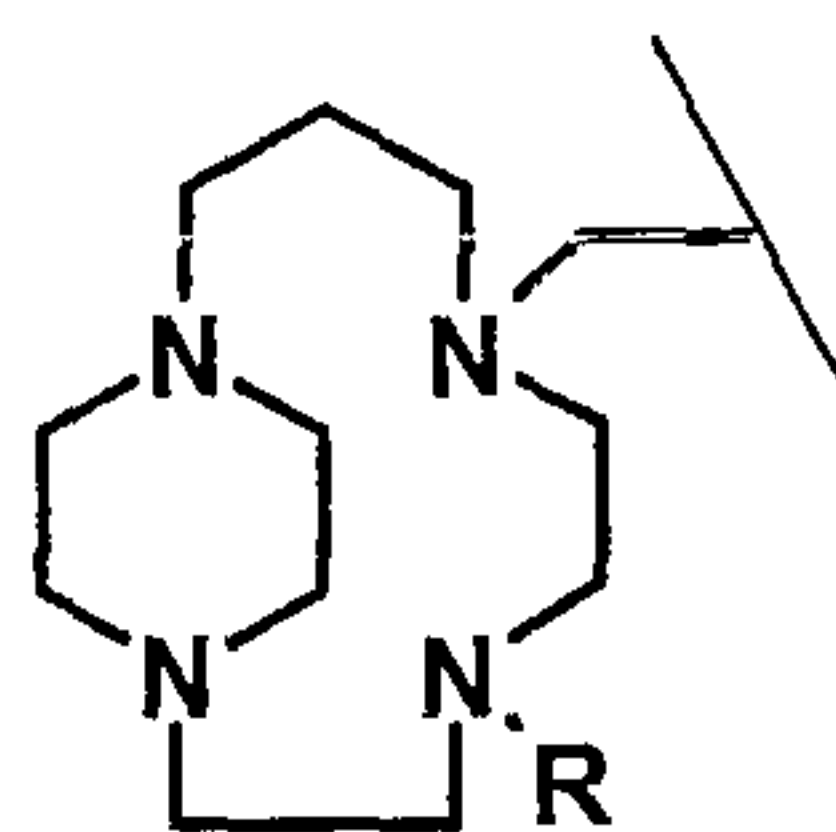
In preferred compounds of Formula 1, 2 or 3, A₁ and A₂ are different. In this
15 case the compound is termed 'unsymmetrical'. In particularly preferred compounds A₁, A₂ and A₃ will be represented by Formulae 7 to 15.



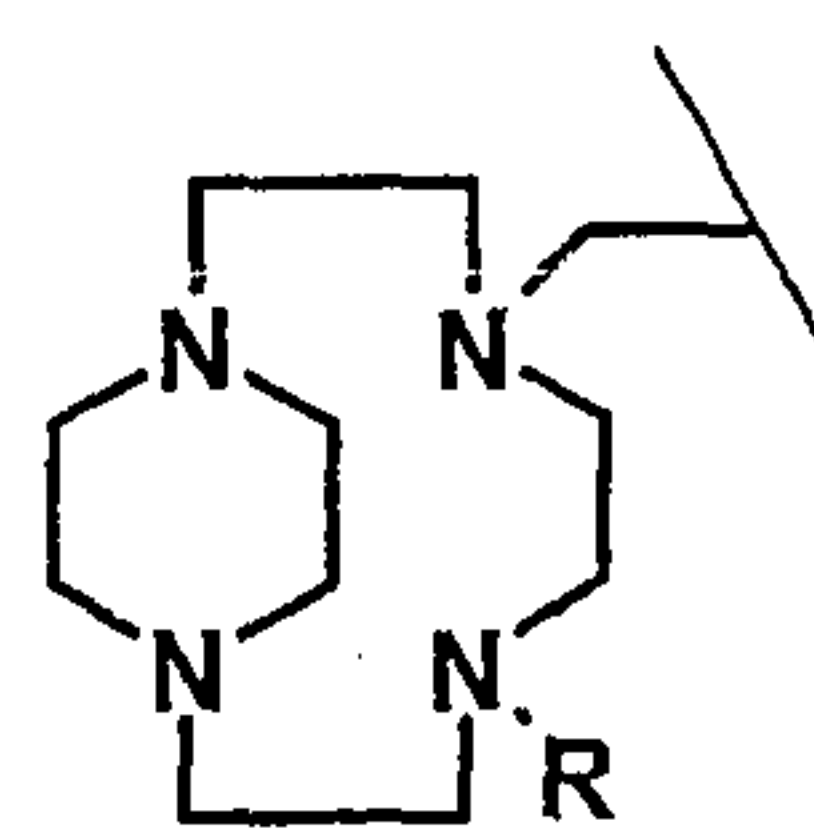
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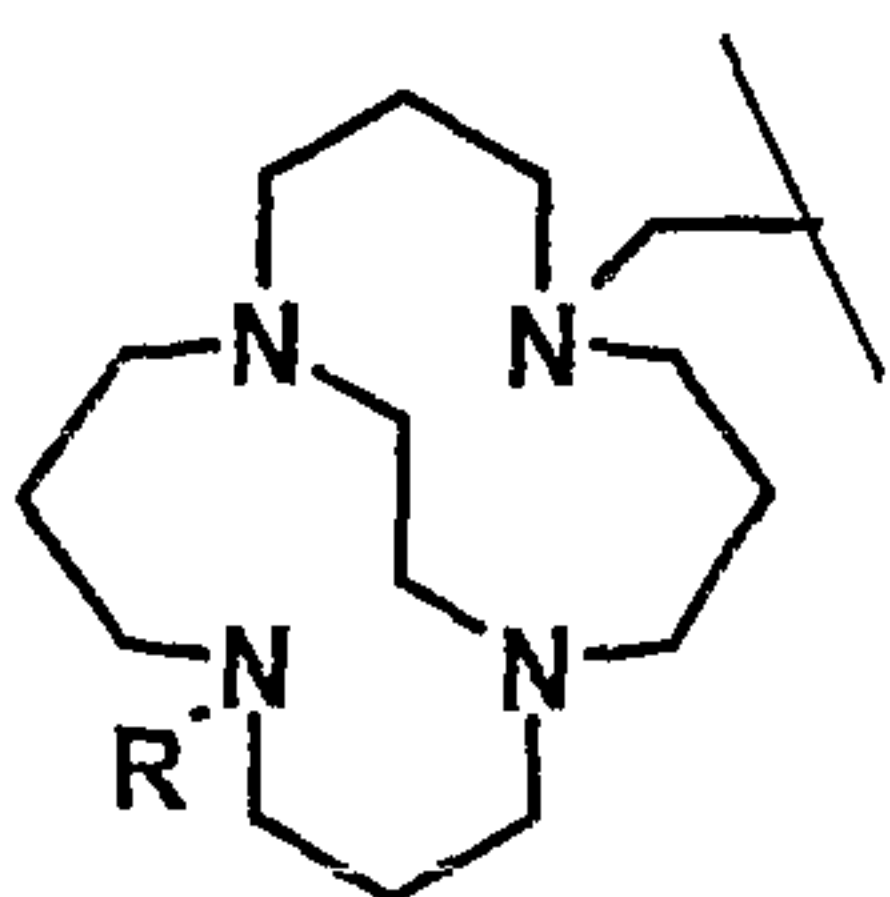
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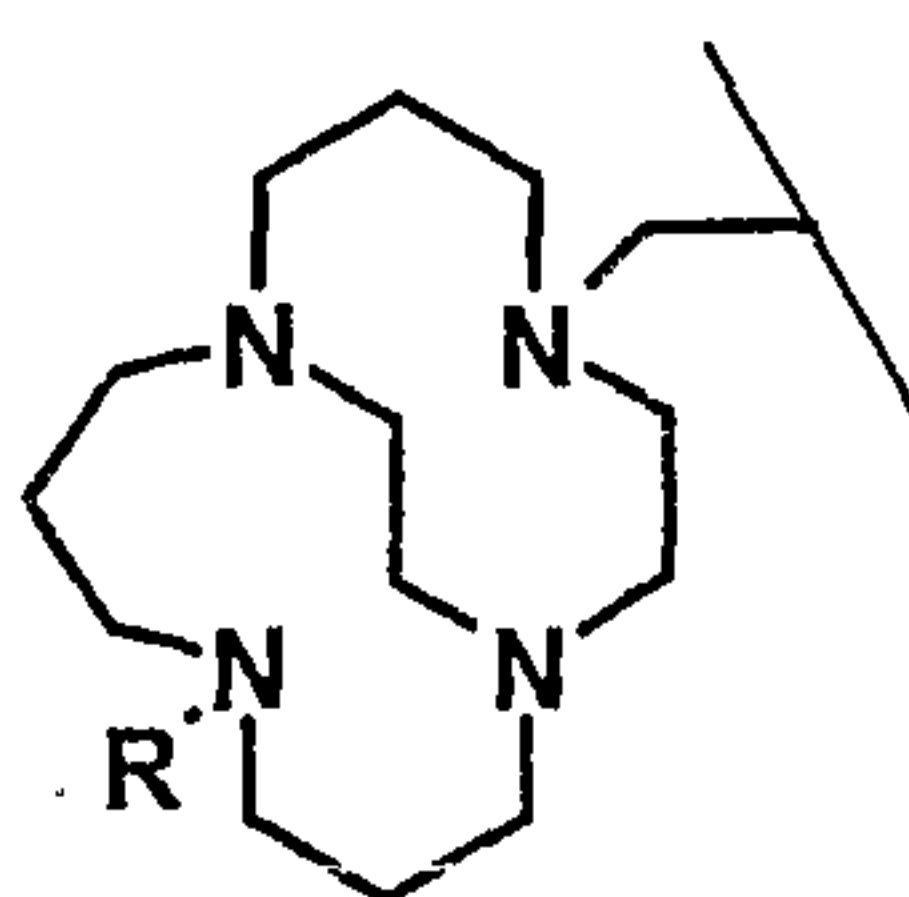
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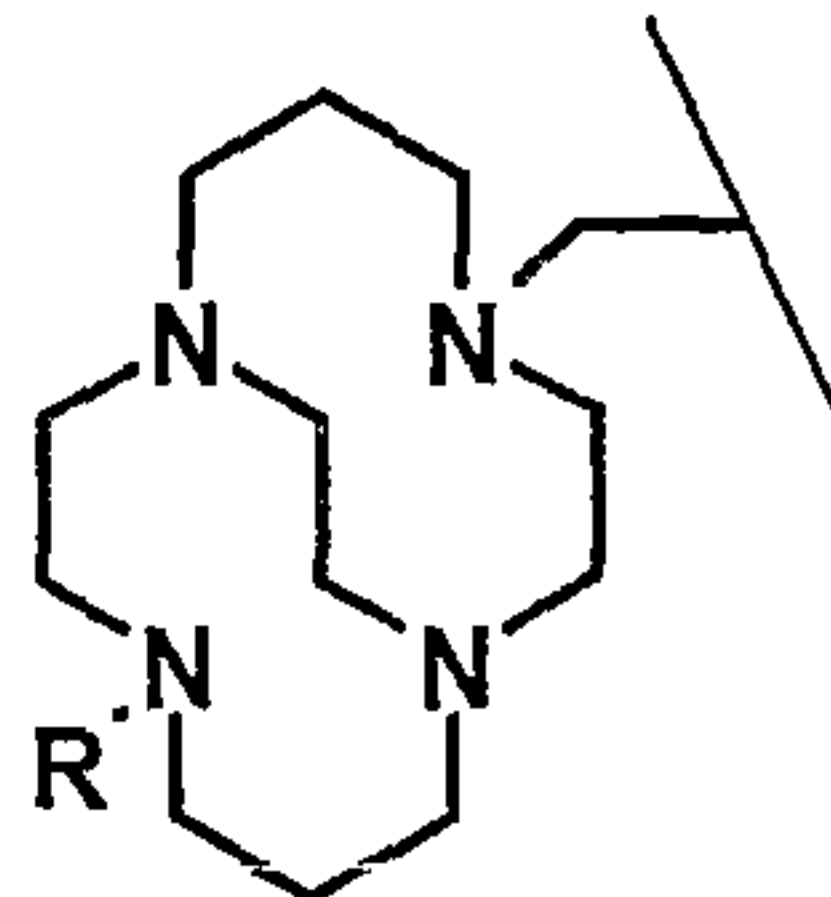
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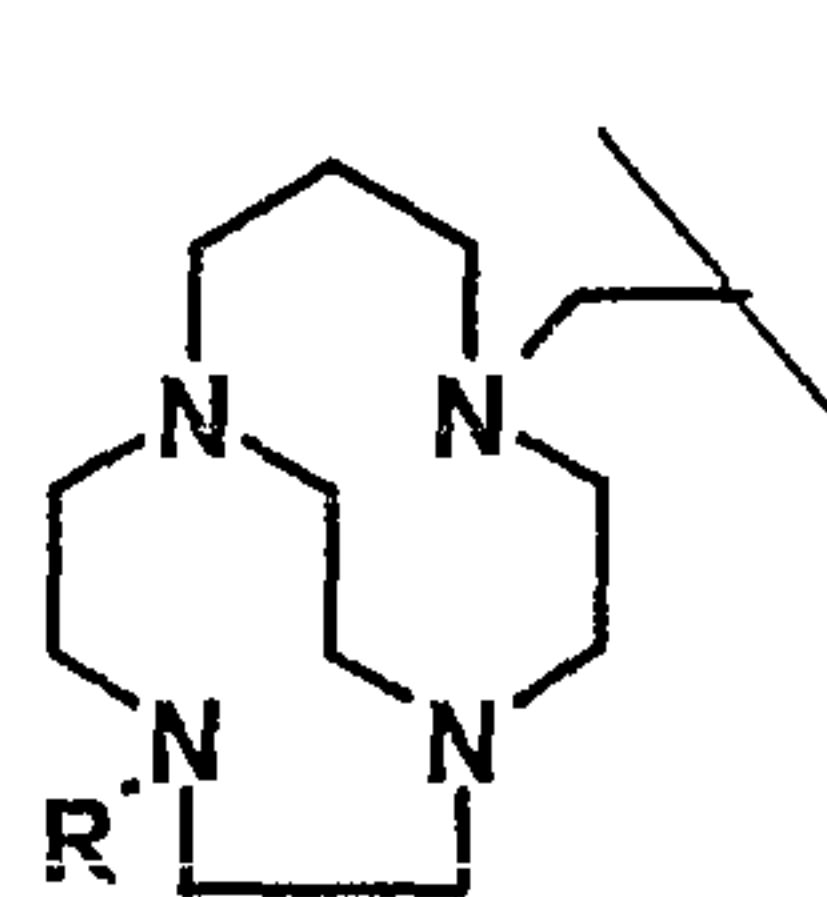
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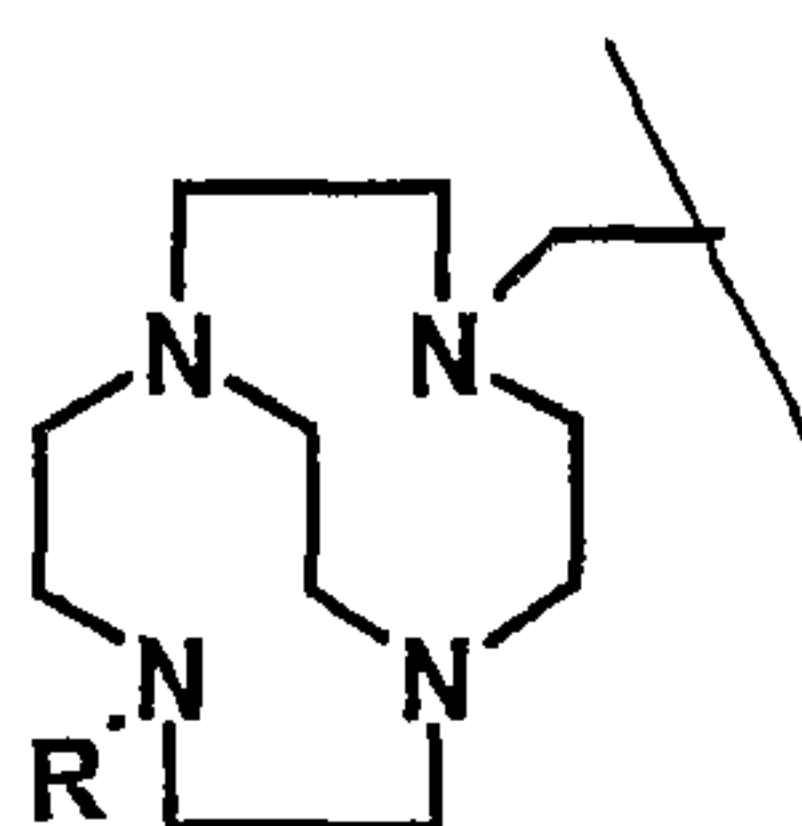
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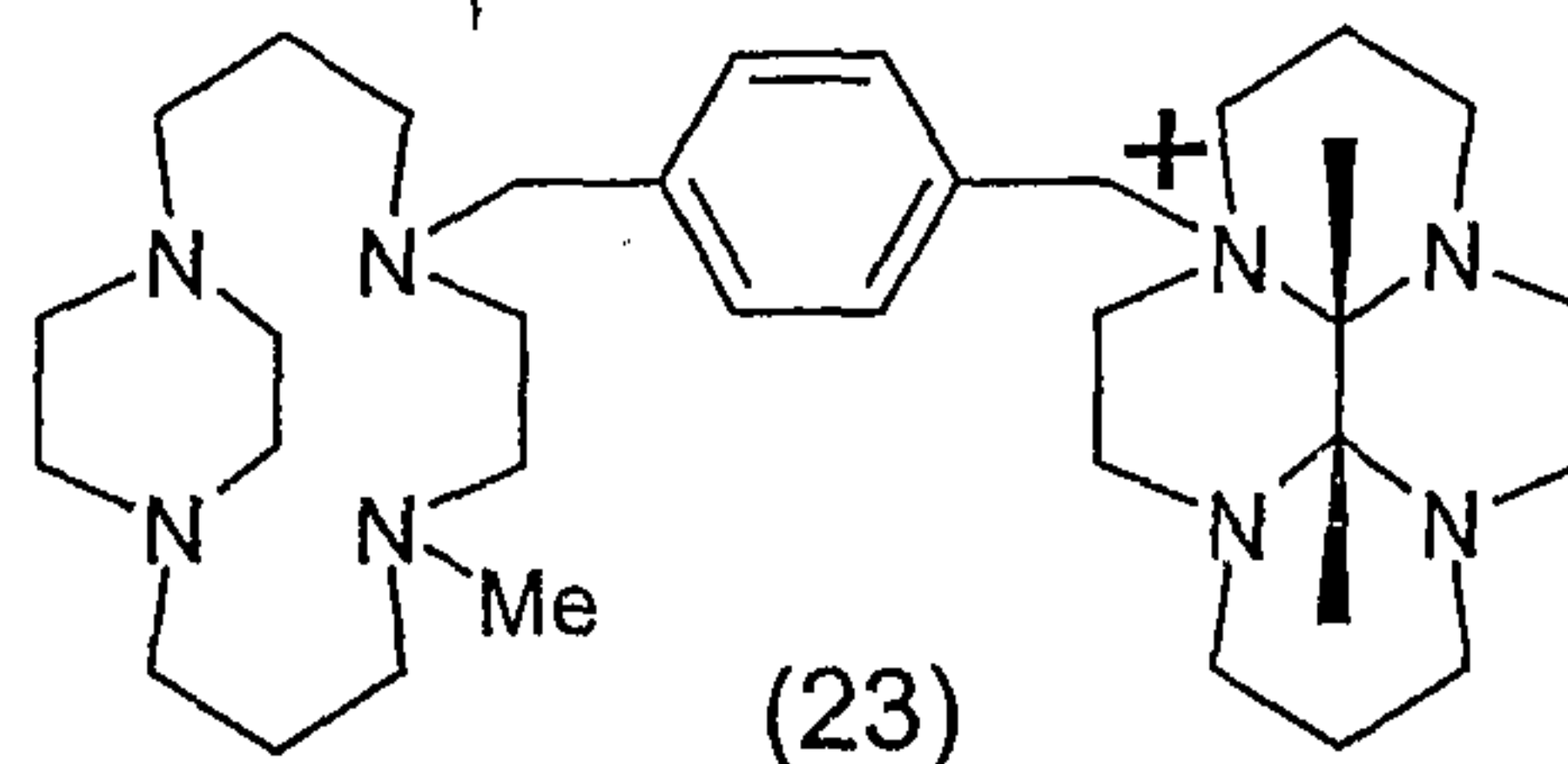
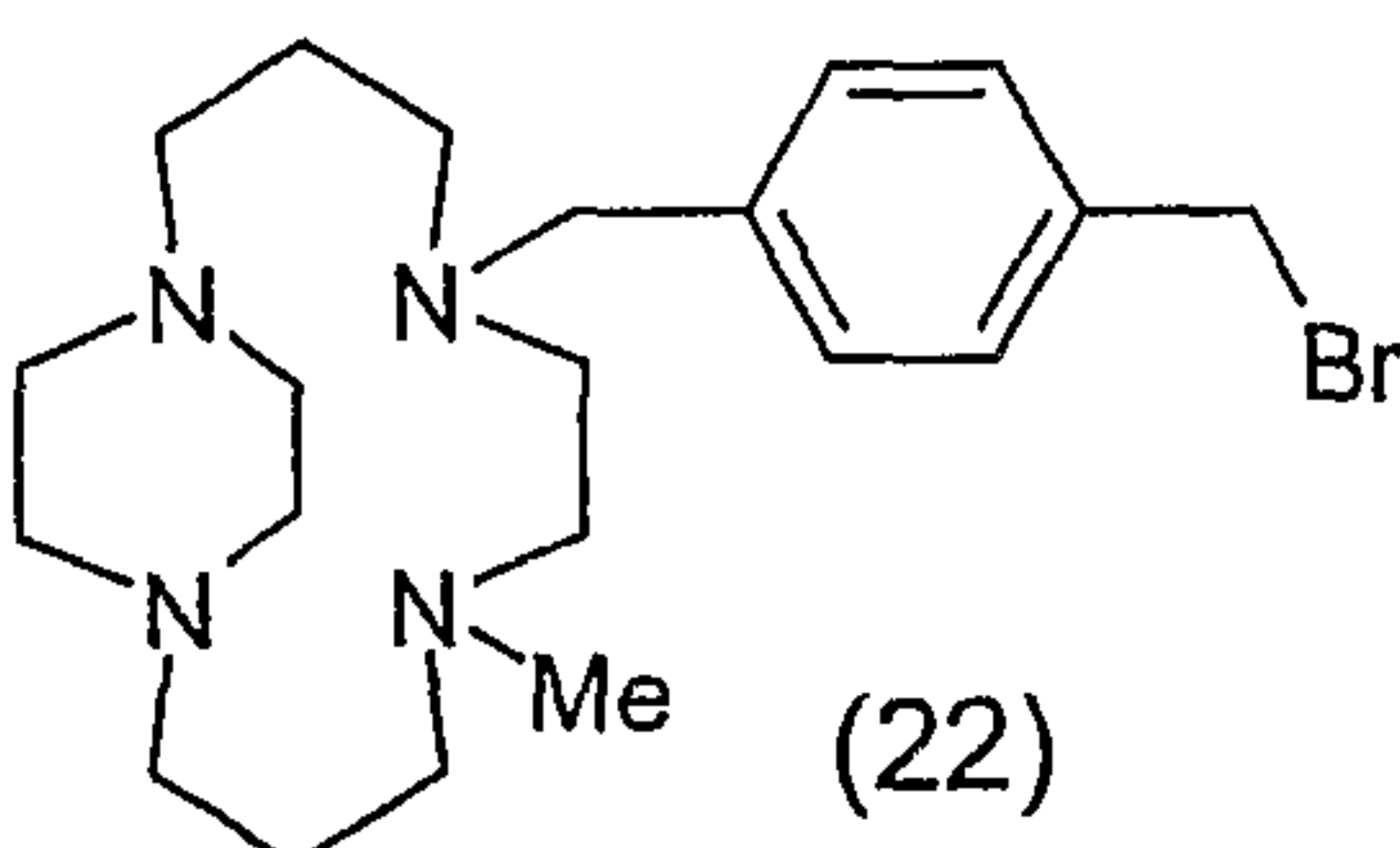
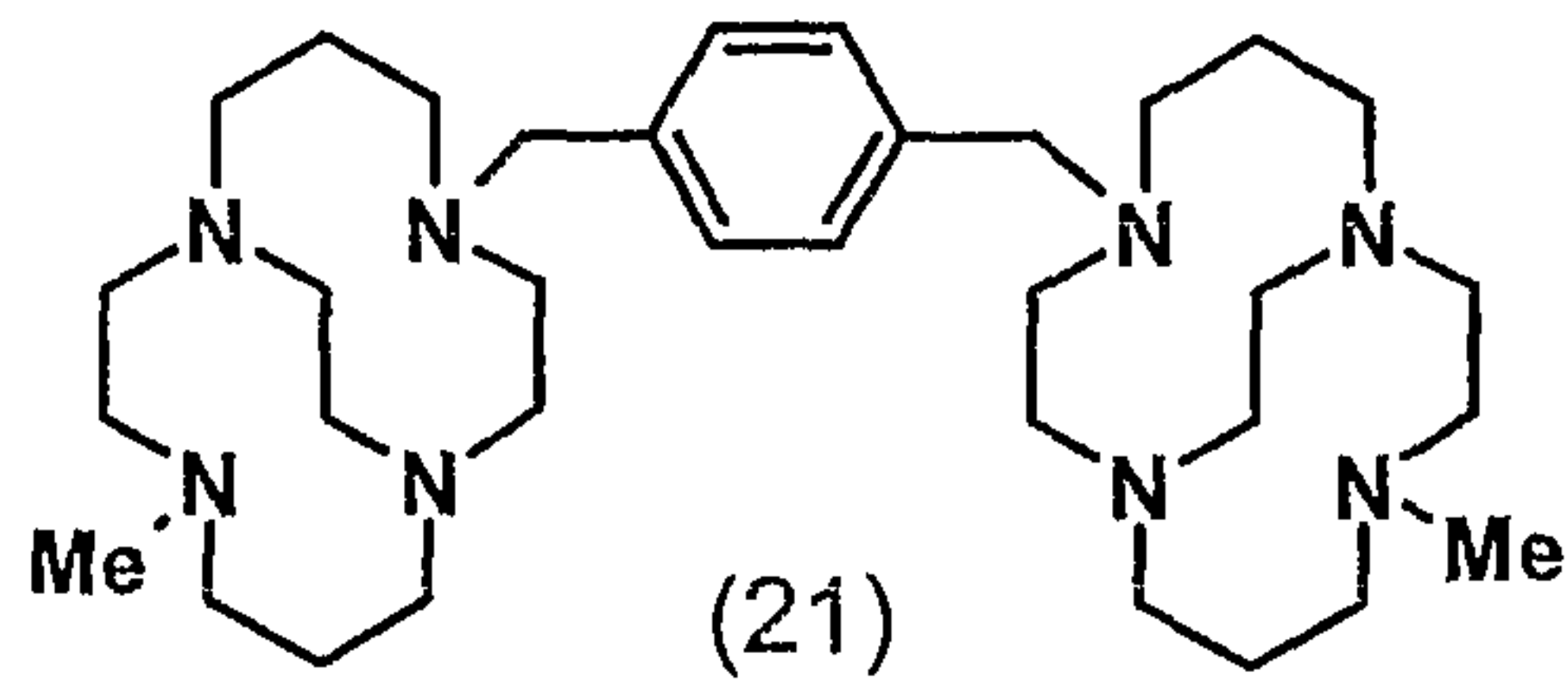
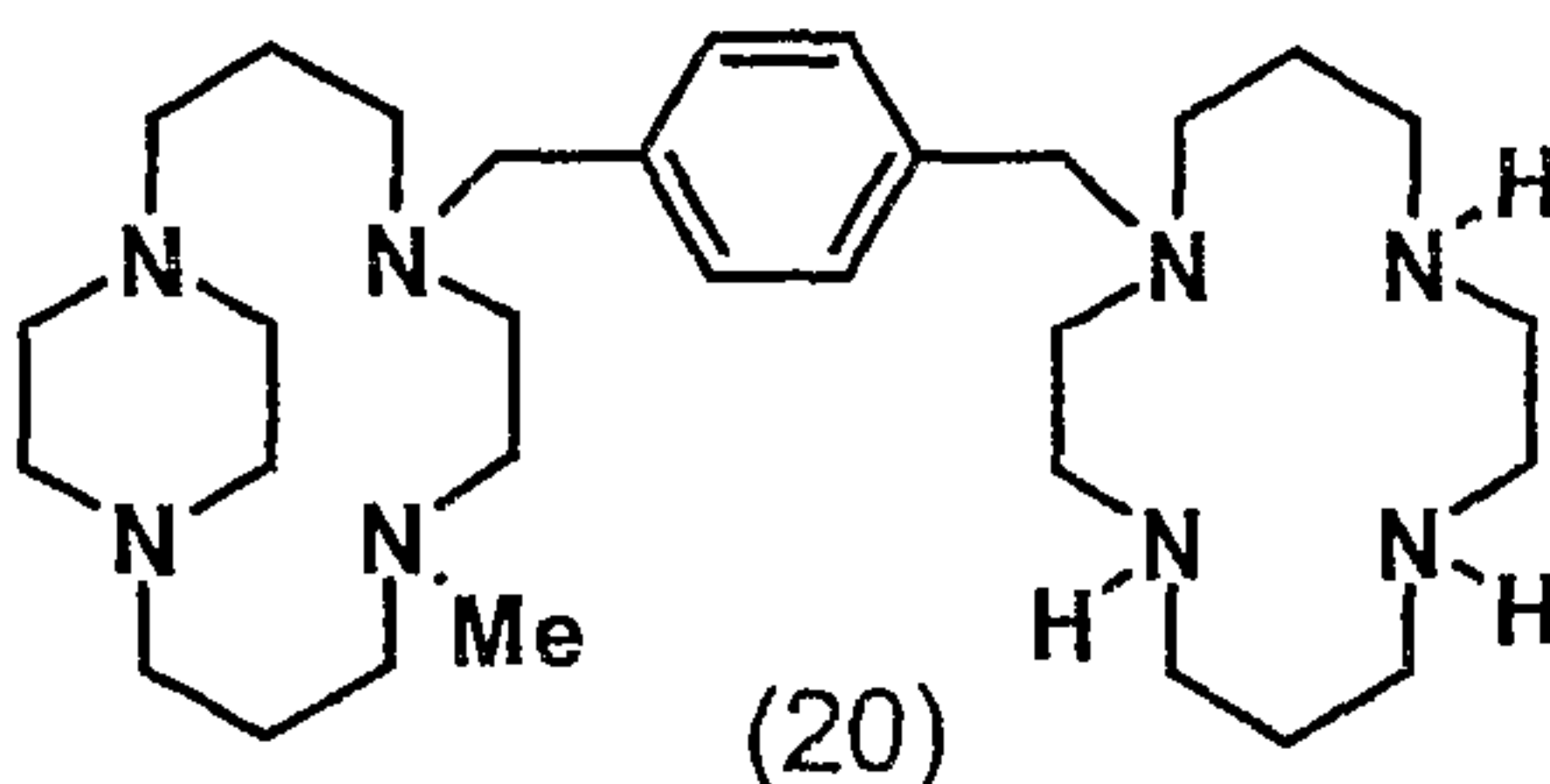
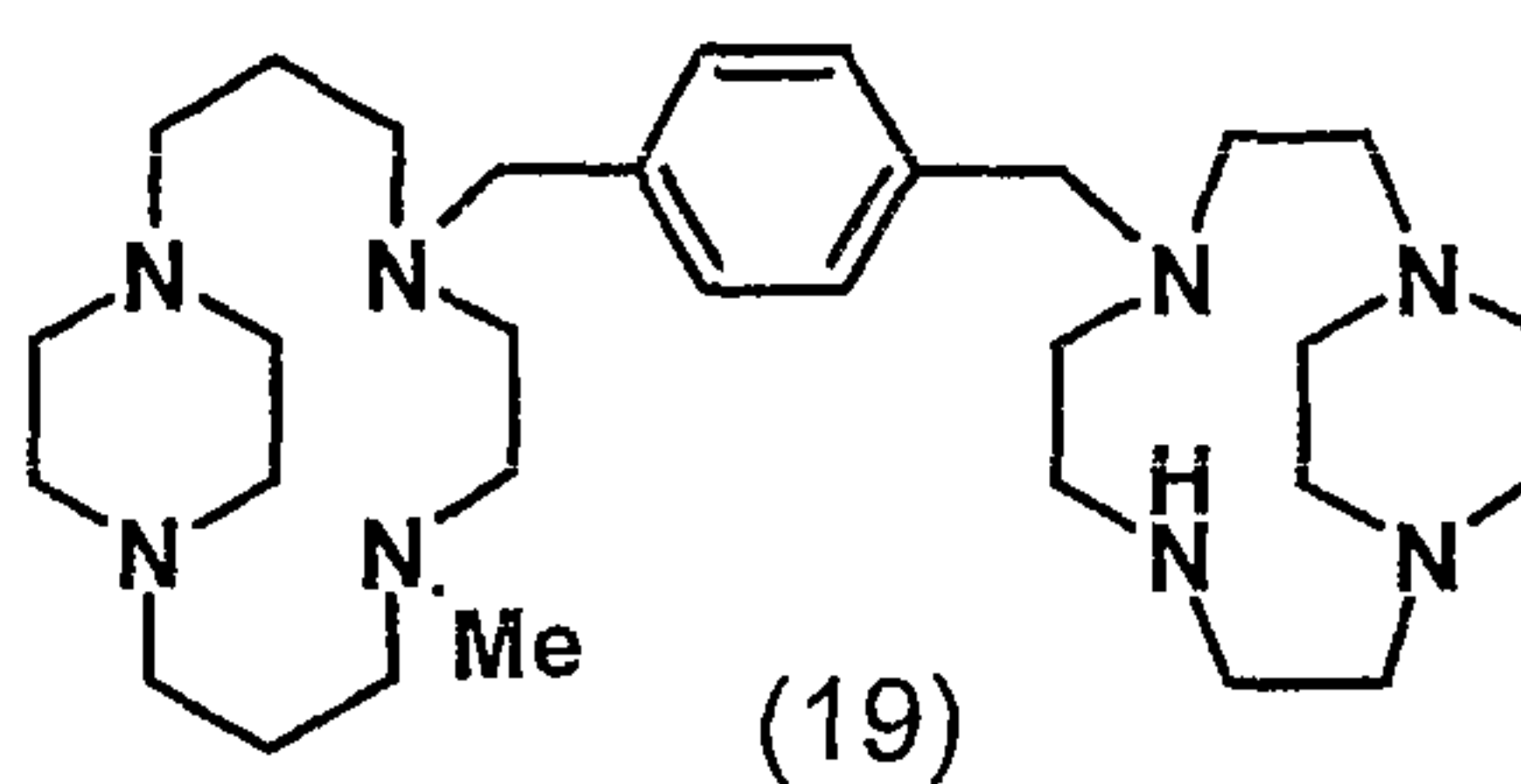
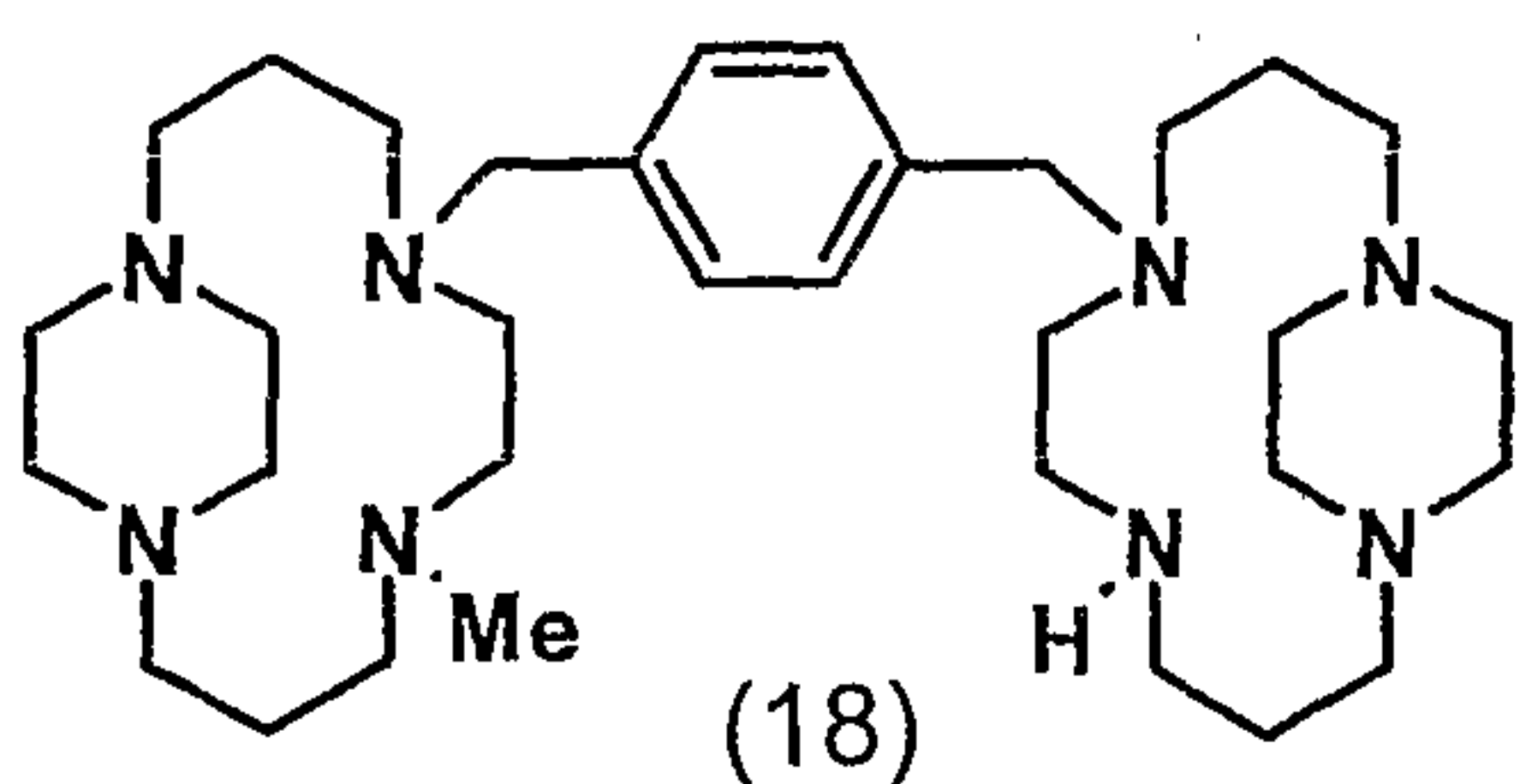
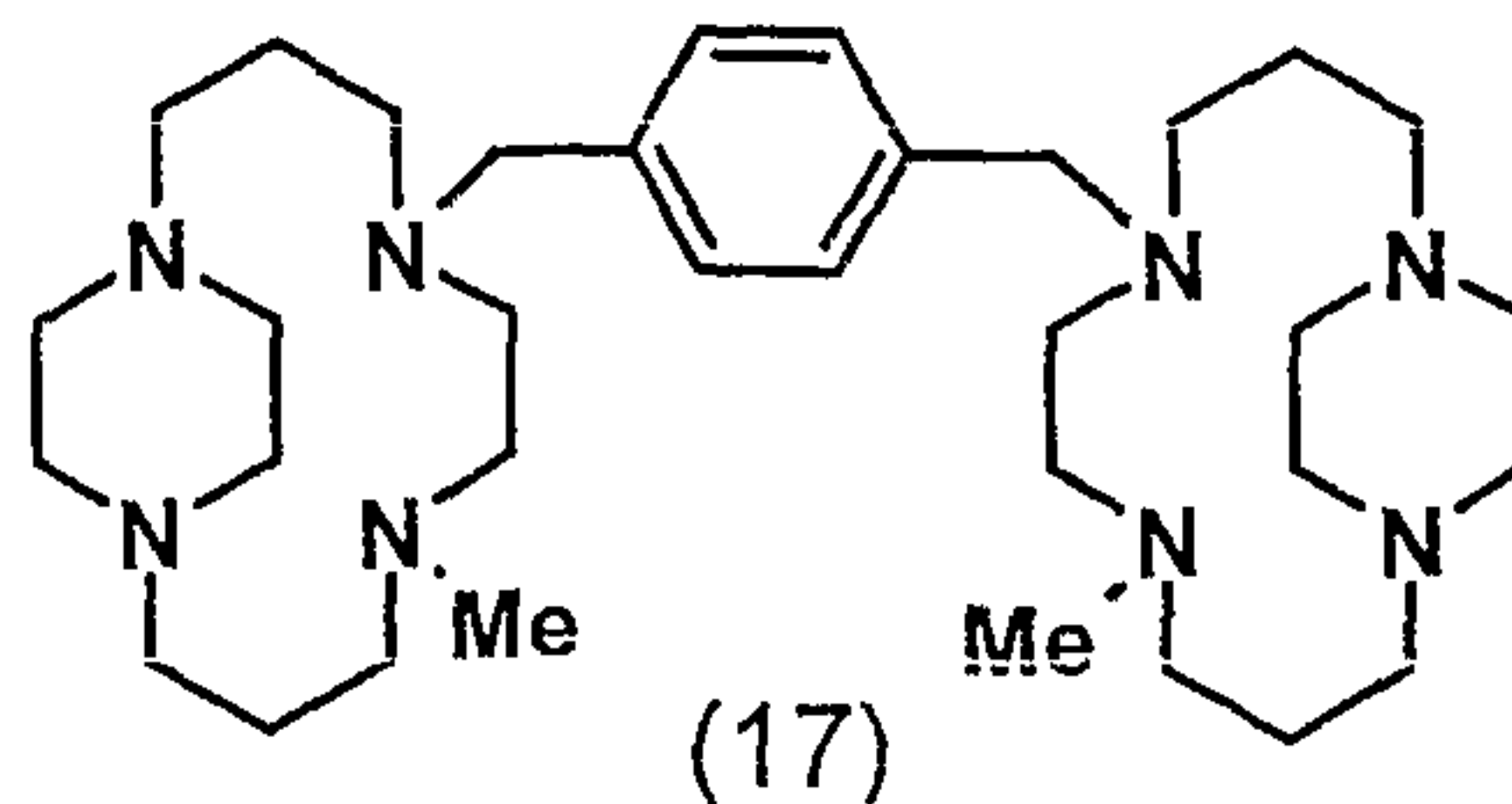
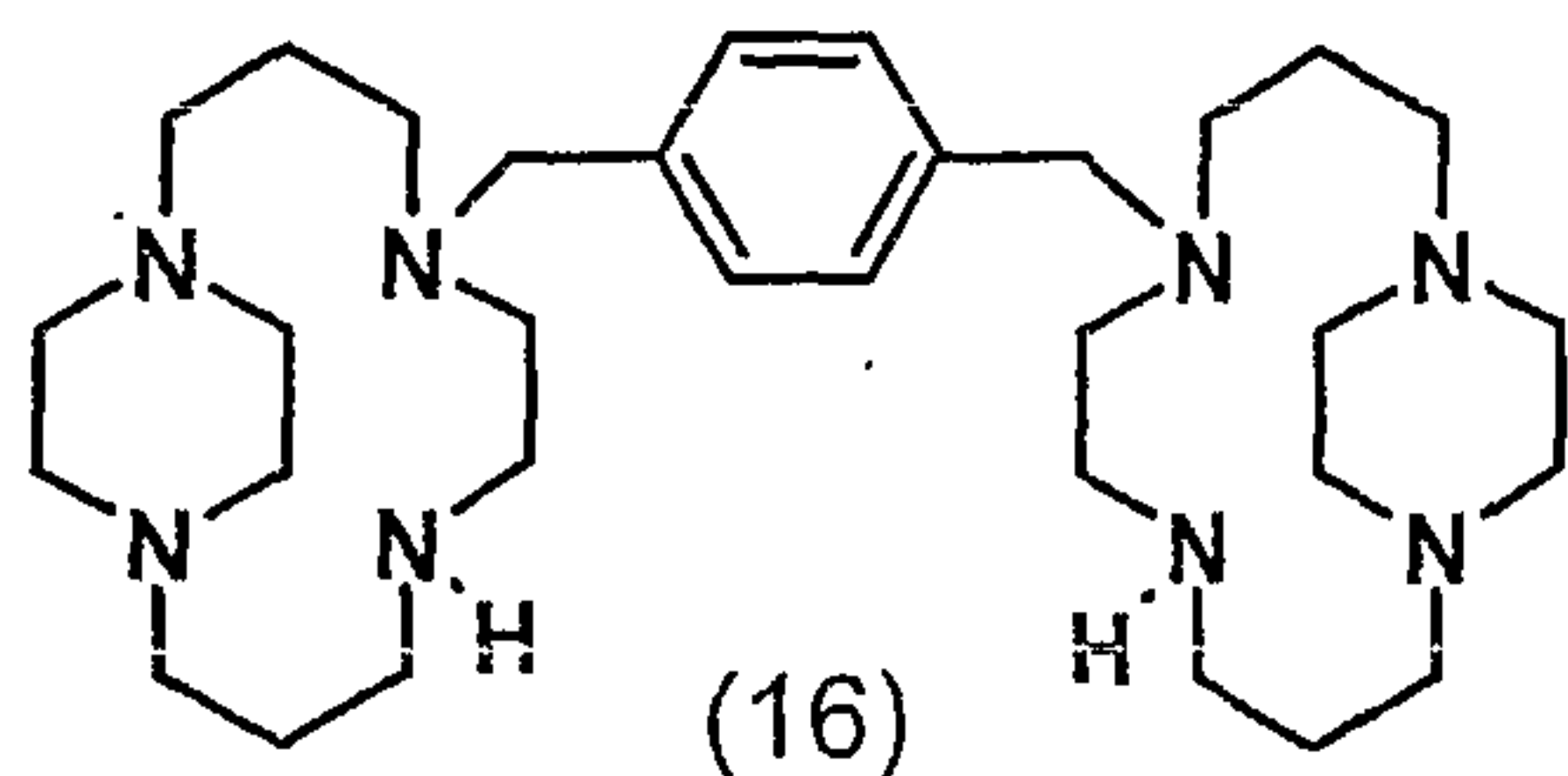


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20

In particularly preferred compounds one or both of R is H or Me.

Preferred compounds may be selected from Formulae 16 to 23:



Preferred compounds of the present invention may bind to CXCR4 chemokine cell surface receptors and find particular but not exclusive application as CXCR4 co-receptor antagonists which are useful in anti-viral treatment, particularly HIV treatment.

In another aspect this invention provides a method for treating a viral disease in a mammal comprising administering to said mammal an effective amount of a compound of Formula 1, 2 or 3, or a pharmaceutically acceptable salt or pro-drug

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thereof. Compounds of the present invention for application for comparative human and of veterinary treatments.

In another aspect this invention provides a pharmaceutical composition
5 comprising a compound of Formula 1, 2 or 3 and a pharmaceutically inert carrier.

In another aspect such compounds may be labelled with radioactive metal isotopes or fluorescent tags allowing in vitro or in vivo imaging of the chemokine receptors for diagnostic or therapeutic use. Such compounds may also be used in the screening of
10 drugs for chemokine receptor binding.

Compounds of the present invention find application in manufacture of imaging agents, for example by covalently binding, chelating or coupling to a fluorophore or radiosotope. A ^{64}Cu labelled compound may be used. A fluorophore or
15 radioisotope containing moiety may be linked to the compound by covalent substitution to the phenyl group, aryl group or to the macrocycle. Such imaging agents may be used for drug screening for either CXCR4 binding molecules or for chemokine receptor imaging from cellular level to diagnostic imaging of whole organisms.

20

The invention is further described by means of example but not in any limitative sense.

Compounds of the present invention are prepared from readily available
25 starting materials.

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to mono- or poly-azamacrocyclic complexes that
30 incorporate a bridged macrocycle. A further aspect of the invention is the transition metal complexes that can be formed wherein said transition metal may have a valence of +1, +2, +3, +4 or +5. In certain aspects of the invention one or more different transition metals may be incorporated. The macrocyclic compounds or the metal

complexes of the present invention are suitable for use as pharmaceuticals that specifically bind to chemokine receptors.

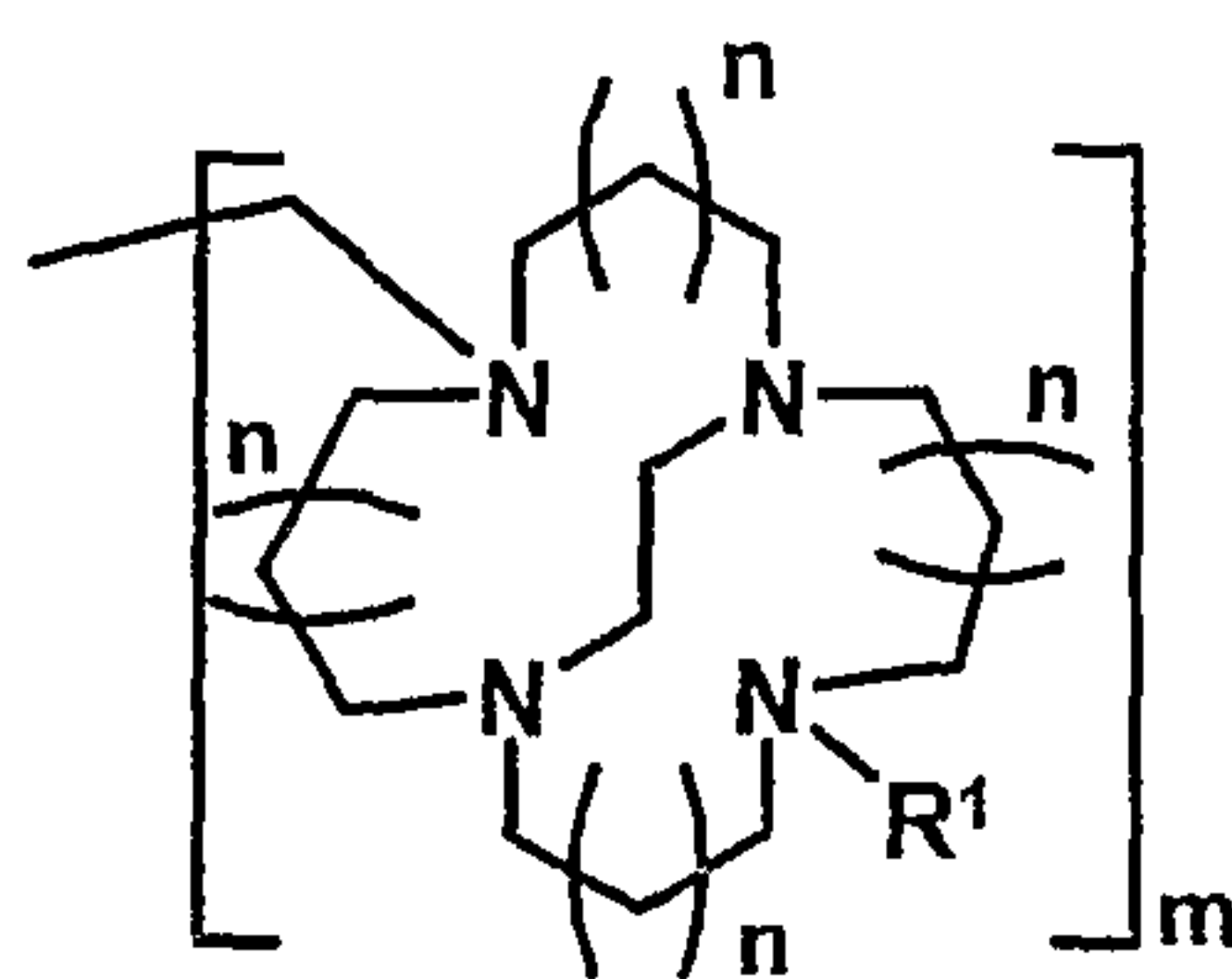
One aspect of the present invention relates to the blocking of a chemokine receptor by binding with a compound of the invention and the subsequent disruption of the chemokine in another role.

In a preferred embodiment one bridged macrocyclic ring (A_1), of the type shown in the non-limiting examples of Formulae 7 to 15, is attached to the aryl ring, Formula 1.

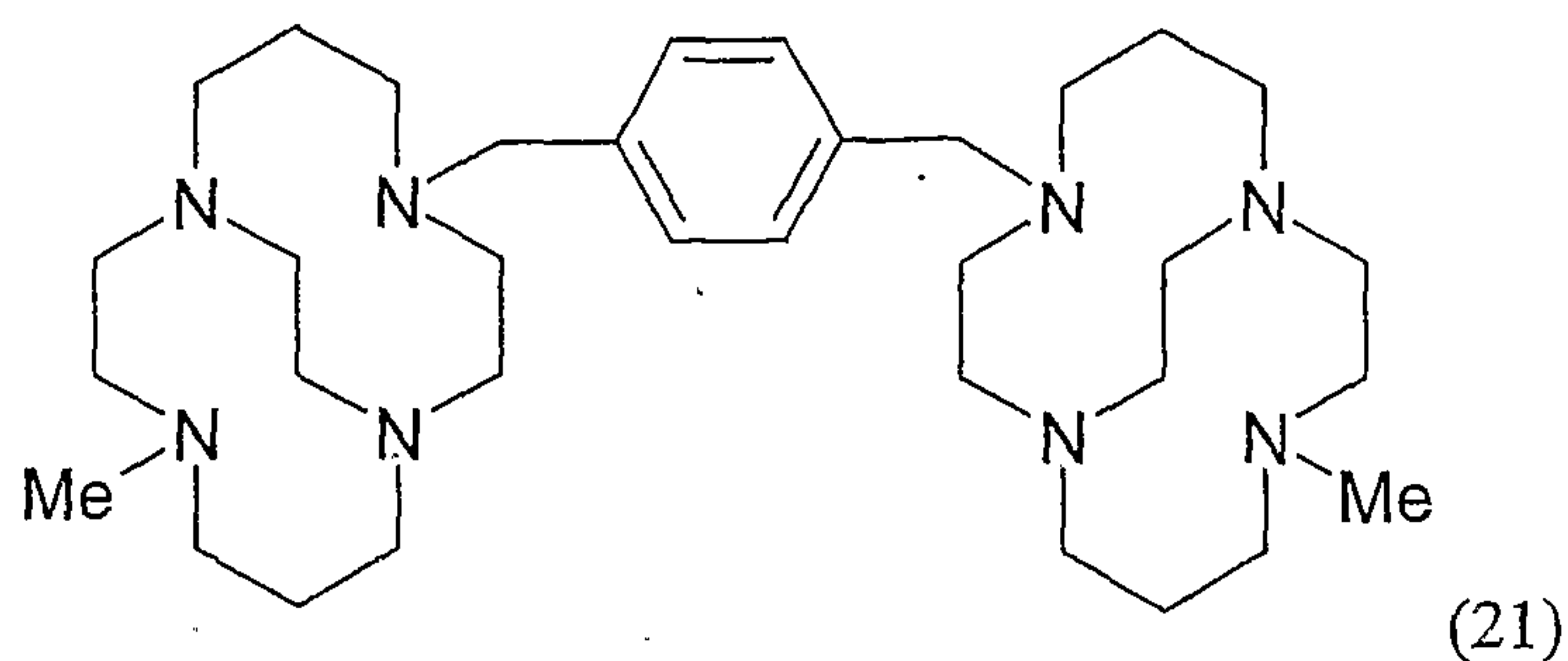
In a further preferred embodiment three macrocyclic rings, with one or more of the type shown in the non-limiting examples of Formulae 7 to 15, are attached to the aryl ring, Formula 3.

In the most preferred embodiment two macrocyclic rings are attached to the aryl ring, with one or more from the non-limiting examples of Formulae 7 to 15, giving the preferred compounds shown of Formula 16 to 23.

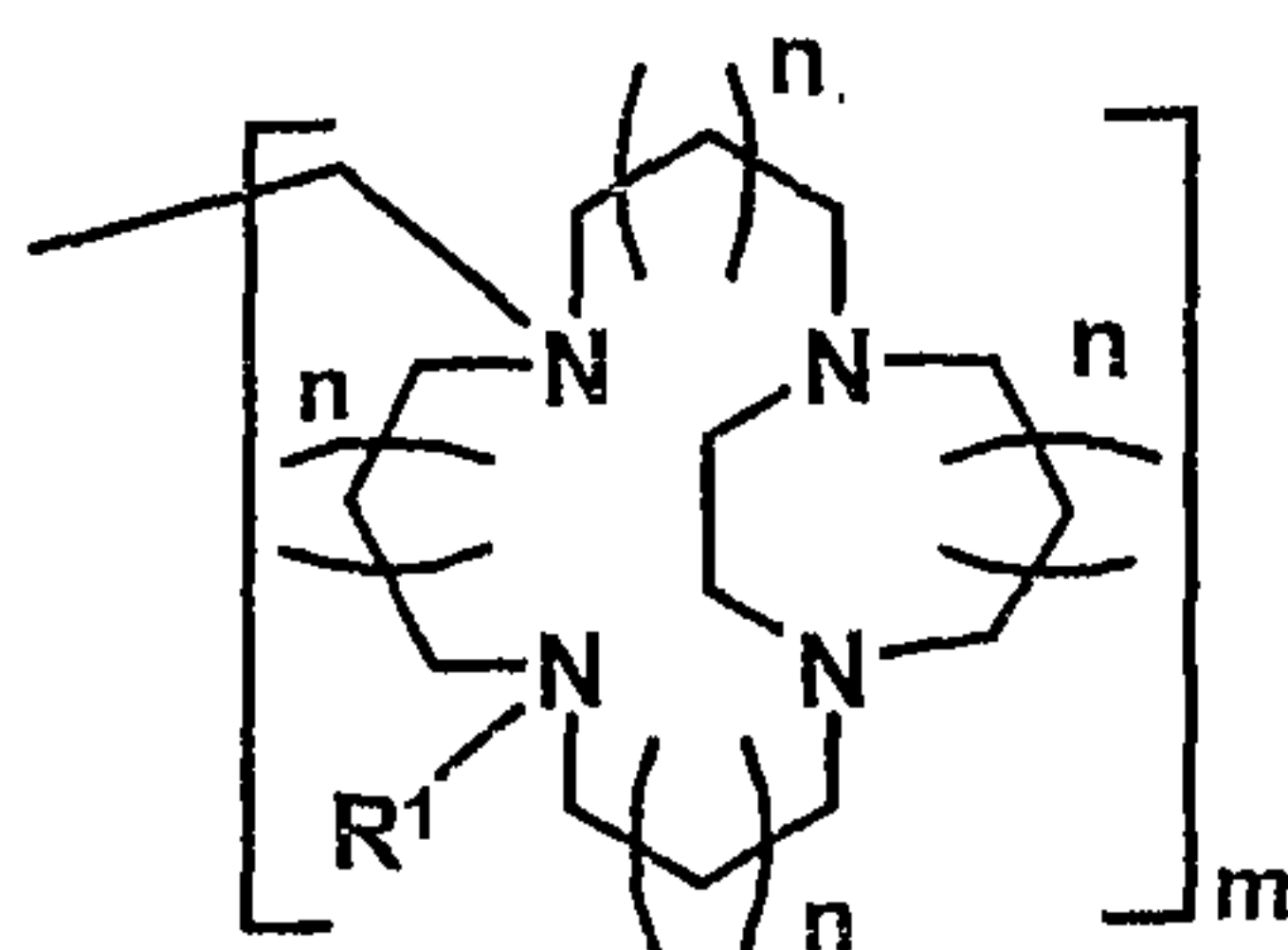
In a preferred embodiment the two carbon linker of at least one macrocycle connects two non-adjacent nitrogens within the ring:



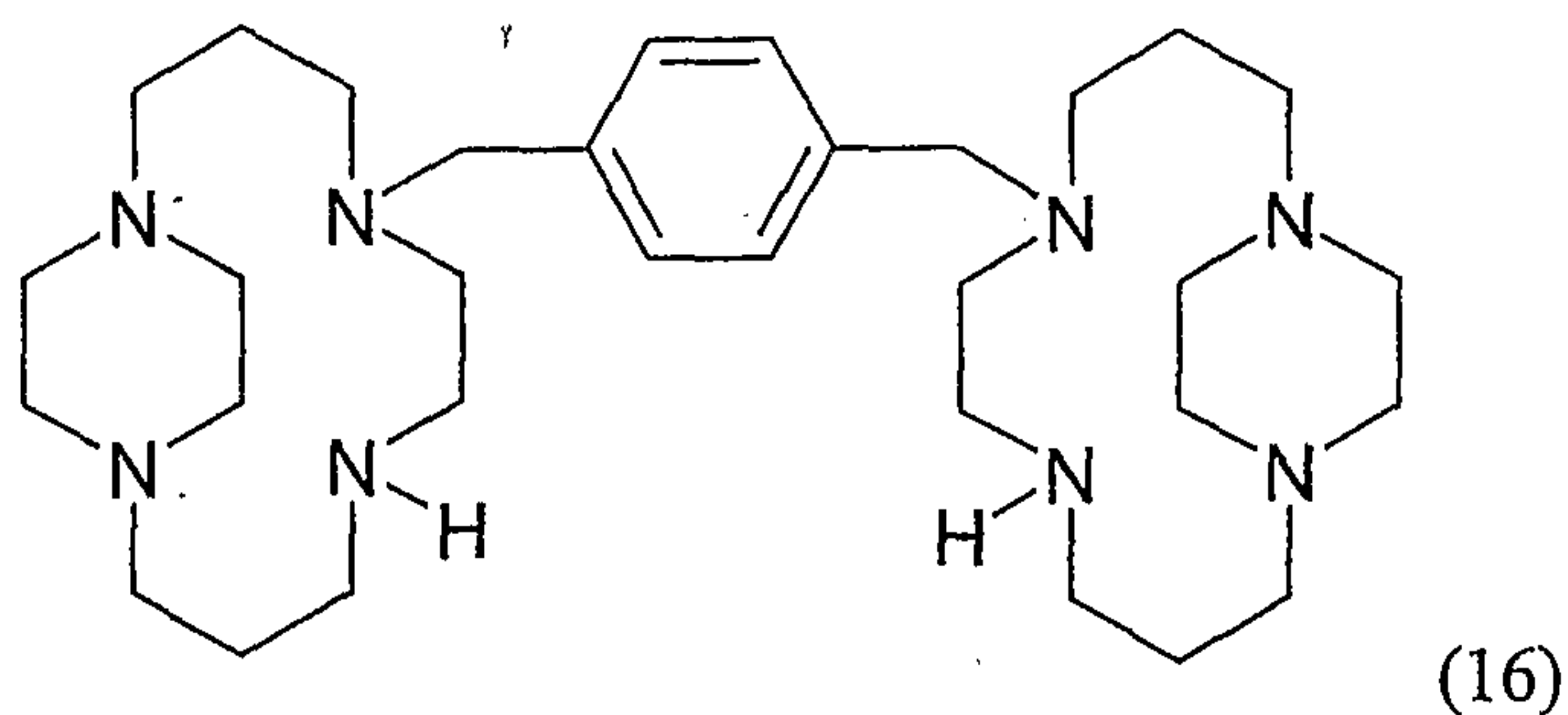
In this embodiment n is 0 or 1 and R is an alkyl group of linear chain length one to six carbons, preferably C_1 to C_3 , with the following structure preferred:



In a preferred embodiment the two carbon linker of at least one macrocycle connects two adjacent nitrogens within the larger ring to form a six membered piperazine ring:
 5 for example with a 1,2-ethyl linker.

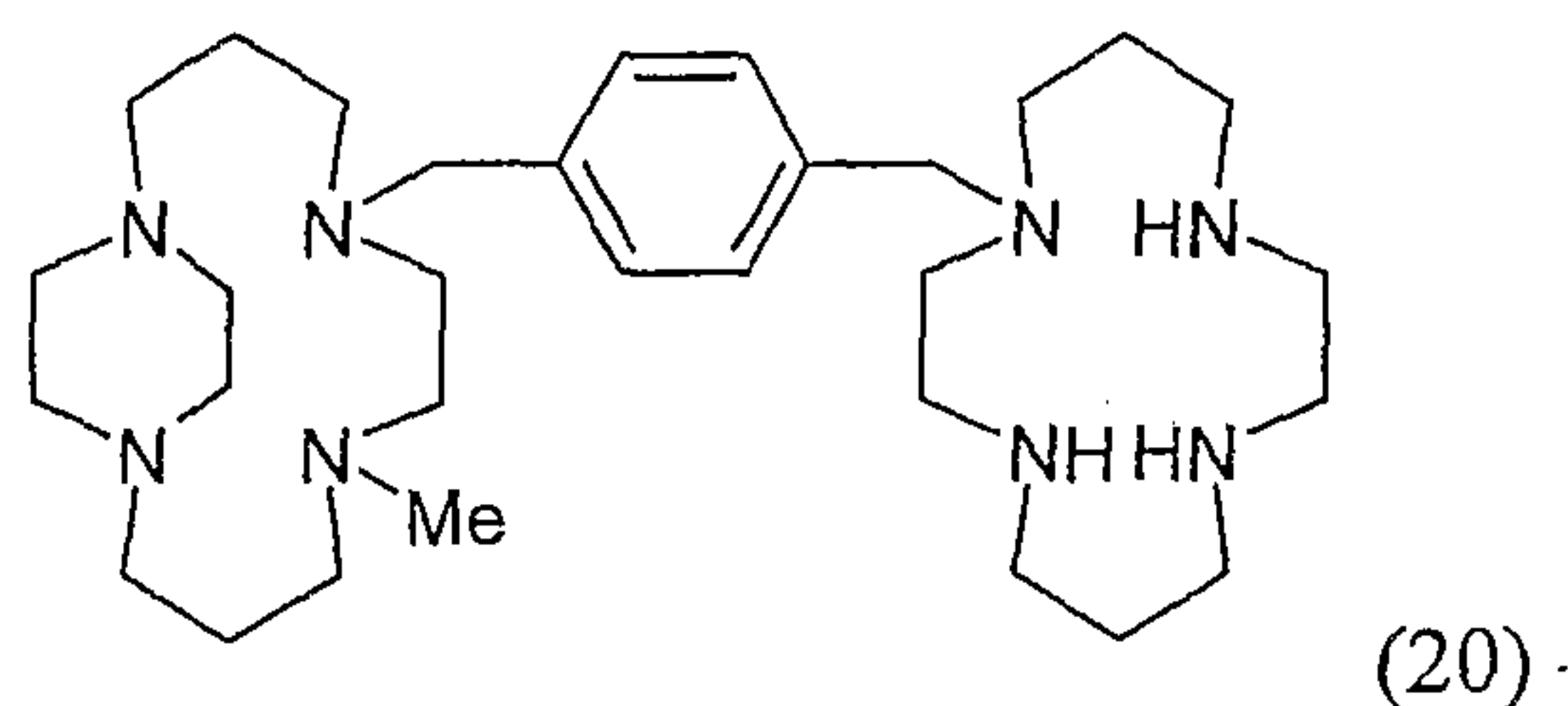


In this embodiment n is an integer from 0 or 1 and R^1 is an alkyl group of linear chain length one to six carbons, preferably C_1 to C_3 , with the following structure preferred:



In a further embodiment there are one or more macrocyclic rings of different type arranged around the aryl centre, Ar (Ar = 1,2-phenyl, 1,3-phenyl, 1,4-phenyl, 1,3,5-phenyl).
 15

In this embodiment the following structure is preferred:



Transition metal complexes

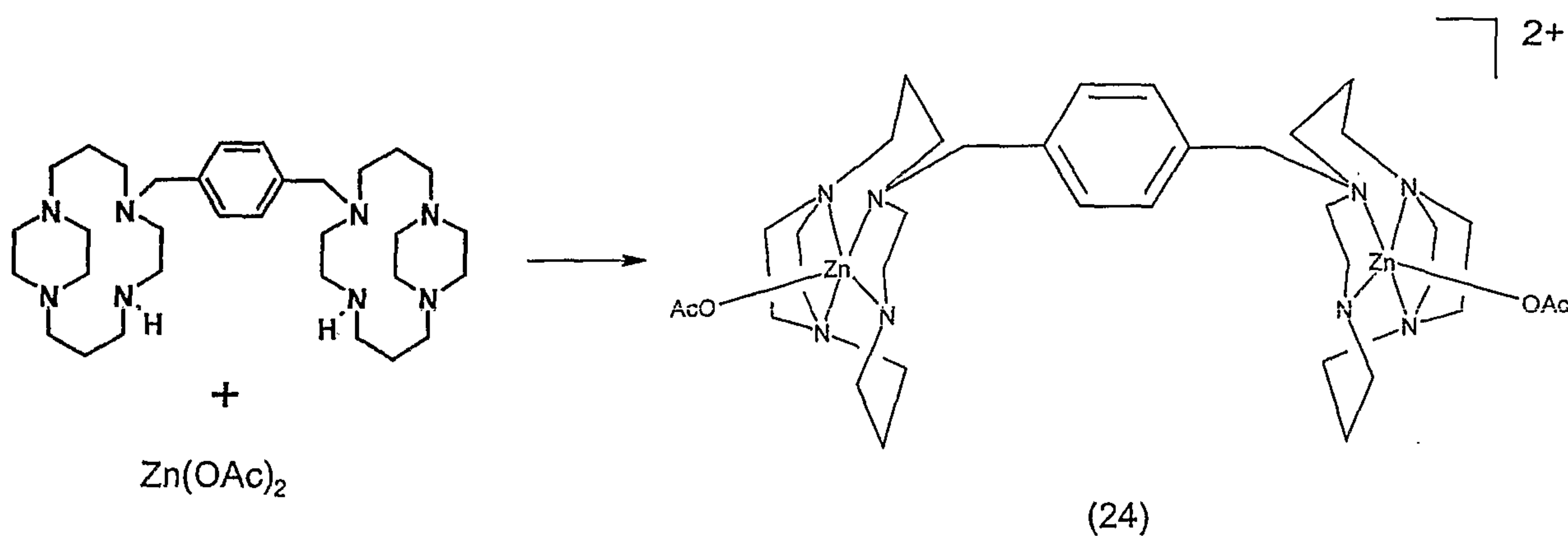
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The transition metal complexes of the present invention comprise one or more transition metals and a ligand with one or more macrocyclic chelators. The complexes of the present invention are charged species having formulae of the type shown in non-limiting examples 24 and 25. The transition metal may have valence of +1, +2, +3, +4 or +5. In these cases a pharmaceutically acceptable anion would be present in sufficient quantity to provide electronic neutrality. Non-limiting examples of preferred transition metals include Fe, Mn, Co, Ni, Cu and Zn.

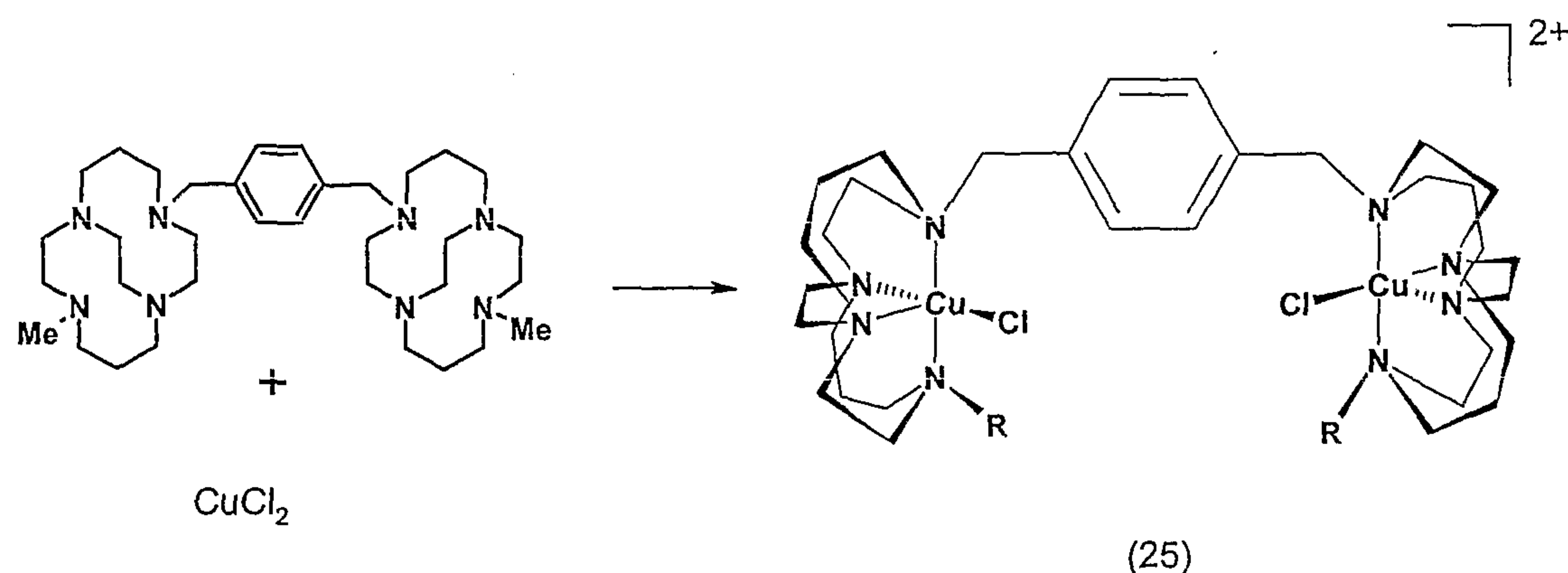
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Preferred examples of the linking aryl unit comprise a 1,4-substituted phenyl unit, a 1,2-substituted phenyl unit, a 1,3-substituted phenyl unit or a 1,3,5-substituted phenyl unit. The most preferred substitution being 1,4-substituted phenyl. The remaining positions of the phenyl unit that are not substituted with macrocycles (A_1 , A_2 or A_3) may be unsubstituted or alkyl substituted (linear alkyl chain C_1 to C_3) with methyl substitution preferred and unsubstituted most preferred.

15



20



Linkage to Fluorophore

Another embodiment of the present invention comprises the attachment of a fluorescent tag to one or more of the macrocyclic chelators either in the presence or absence of one or more transition metal ions. A non-limiting example of a pharmaceutically viable fluorescent tag is Rhodamine B.

Linkage to Radioisotope

Another embodiment of the present invention comprises of the insertion of one or more radioisotopes into one or more of the macrocyclic chelators either in the presence or absence of one or more transition metal ions. A non-limiting example of a pharmaceutically significant radioisotope for imaging applications is ^{64}Cu [following a similar procedure to that described by X. Sun *et al.* Journal of Medicinal Chemistry, 2002, 45, 469.]

Other aspects

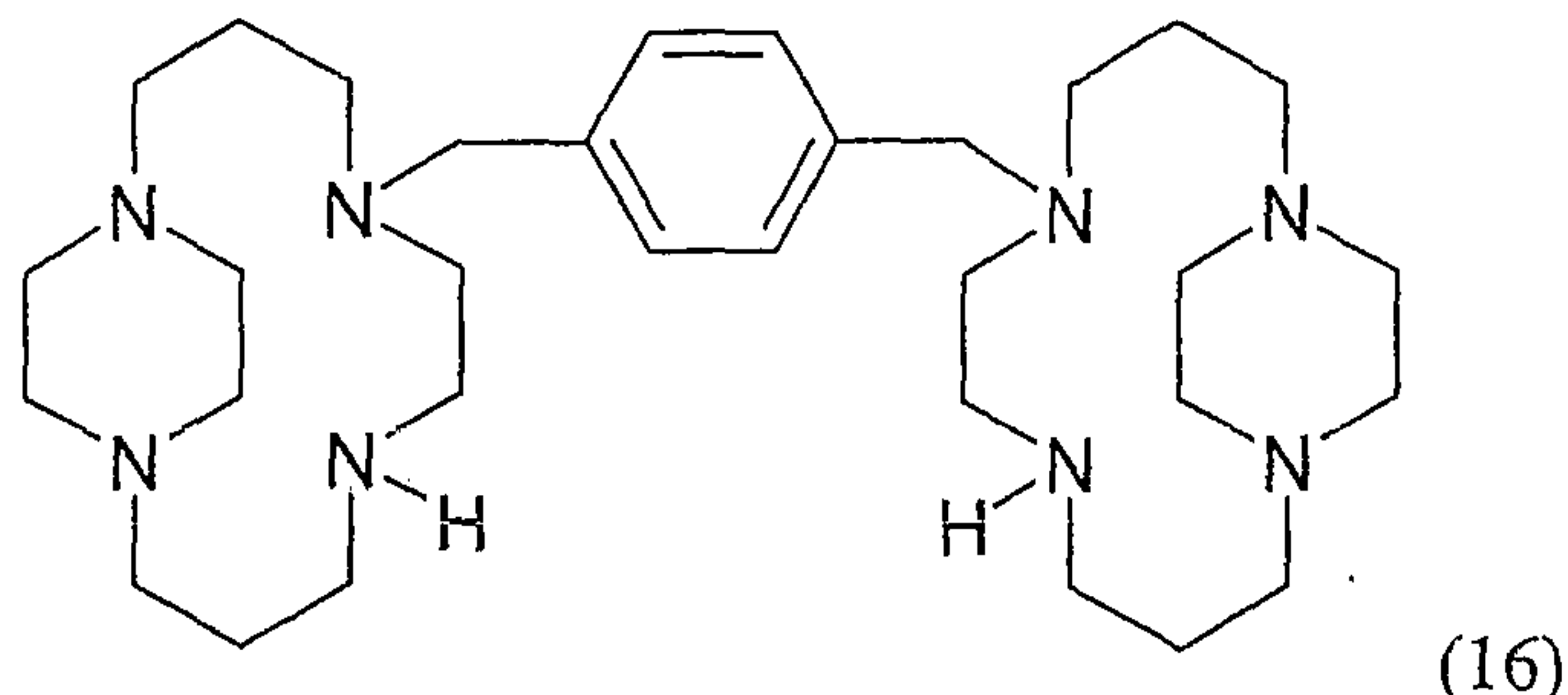
Another embodiment of the present invention relates to a method of treating viral disorders which comprises administering to a mammal an amount of a compound selected from Formulae 1, 2 or 3, effective for treating the disorder, optionally together with one or more excipients or pharmaceutical carriers.

Generally, a sterile aqueous solution or a physiological or buffered solution of the invention or of its metal complexes will be administered to the patient in a variety of ways, including orally, intrathecally and especially intravenously. The invention may also be co-administered with other pharmaceuticals for anti-viral treatment.

EXAMPLES

Example 1 *para*-Xylyl bis(1,5,8,12-tetraazabicyclo[10.2.2]hexadecane

(Compound of Formula 16)



cis-Perhydrotetraazapyrene, 1.2 g (5.40 mmol) and *para*-xylenedibromide, 0.71 g (2.70 mmol) were stirred together in dry acetonitrile (20 ml) for 24 hr. A white precipitate resulted which was collected by filtration, washed in acetonitrile and dried in vacuo to give a white solid (1.35 g, 71%). [Procedure modified from M. Le Baccon *et al*, New Journal of Chemistry, 2001, 25, 118.] Mass and NMR (^1H and ^{13}C) spectra were consistent with the expected product.

This compound, 1.16g (1.60 mmol), was dissolved in dry ethanol (100ml), stirred under nitrogen and NaBH_4 , 0.61g (16 mmol), added over a period of 20 mins. The solution was then stirred for a further 30 mins and refluxed under nitrogen for one hour. The solution was cooled to room temperature, 3M HCl (10ml) was added slowly and the solvent removed. The resulting residue was dissolved in water (30ml) and the pH was reduced to 14 using KOH pellets. Benzene was used to extract the basic solution (6x50ml). The combined extracts were dried, filtered and the solvent removed giving a white solid, (0.59g, 65%):

^1H NMR (400 MHz, CDCl_3) δ 7.20 (s, 4H, ArH), 3.65 (s, 4H, NCH_2Ar), 3.27 (m, 4H, NCH_2), 3.02 (m, 4H, NCH_2), 2.93(m, 4H, NCH_2), 2.70(m, 4H, NCH_2), 2.63(m, 8H, NCH_2), 2.56(m, 12H, NCH_2), 2.26(m, 4H, NCH_2), 1.81 (s, 4H, NCH_2), 1.72 (m, 4H, NCH_2).

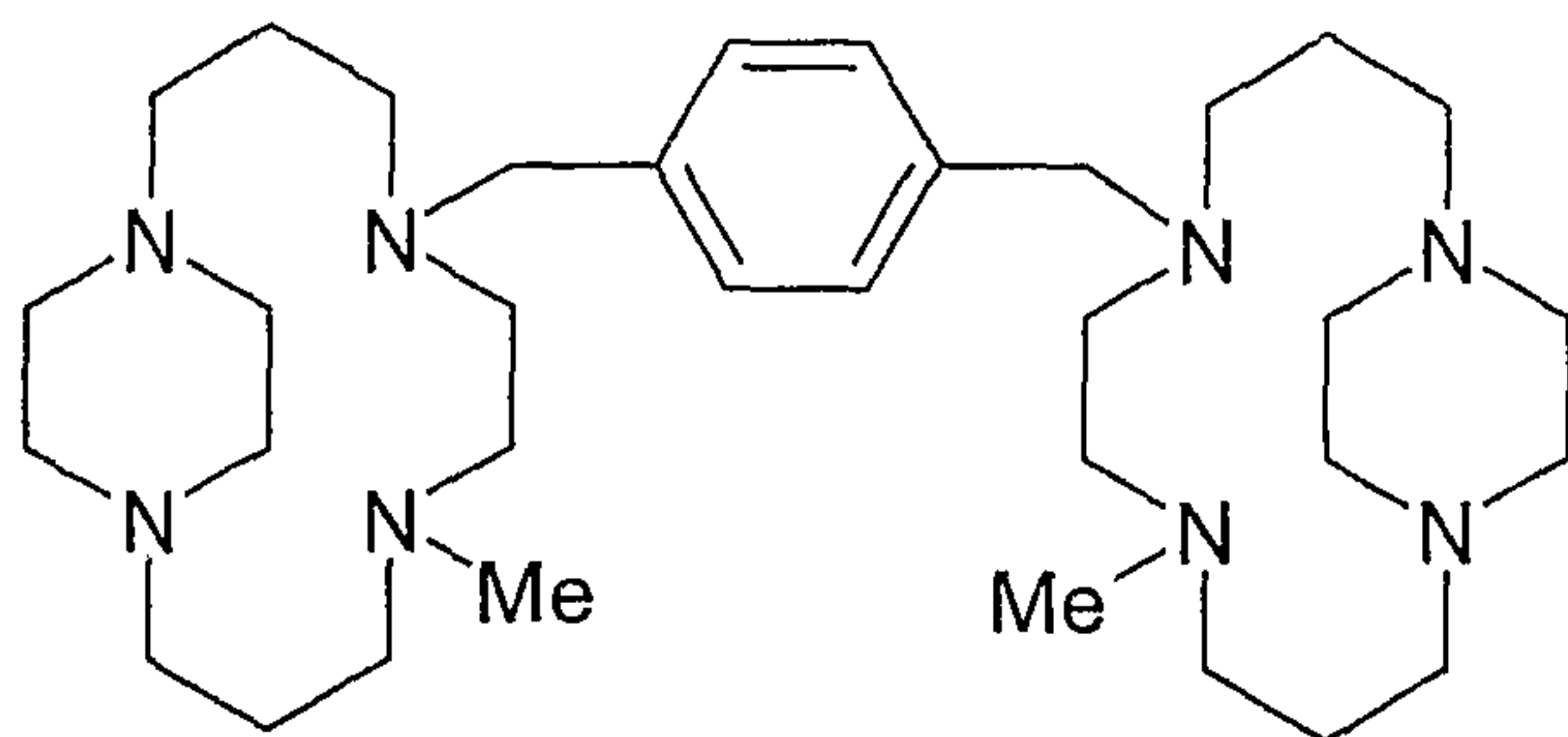
^{13}C NMR (CDCl_3) δ 136.40(*ArCH*₂), 129.36(*ArH*), 57.07(*ArCH*₂N), 56.24(*CH*₂N), 55.51(*CH*₂N), 55.15(*CH*₂N), 54.73(*CH*₂N), 51.30(*CH*₂N), 50.62(*CH*₂N), 48.26(*CH*₂N), 48.17(*CH*₂N), 26.43(*NCH*₂*CH*₂), 23.52(*NCH*₂*CH*₂).

The HCl salt of this compound was formed by following method. The solid, 101mg (0.19mmol), was dissolved in methanol (10ml) and conc HCl:methanol

(1ml:20ml) was added dropwise until a precipitate was observed. Filtrate was decanted off and this process was repeated twice. Finally, the filtrate was concentrated in vacuo to give a white powder as product. Mass and NMR (^1H and ^{13}C) spectra were consistent with the expected product.

5

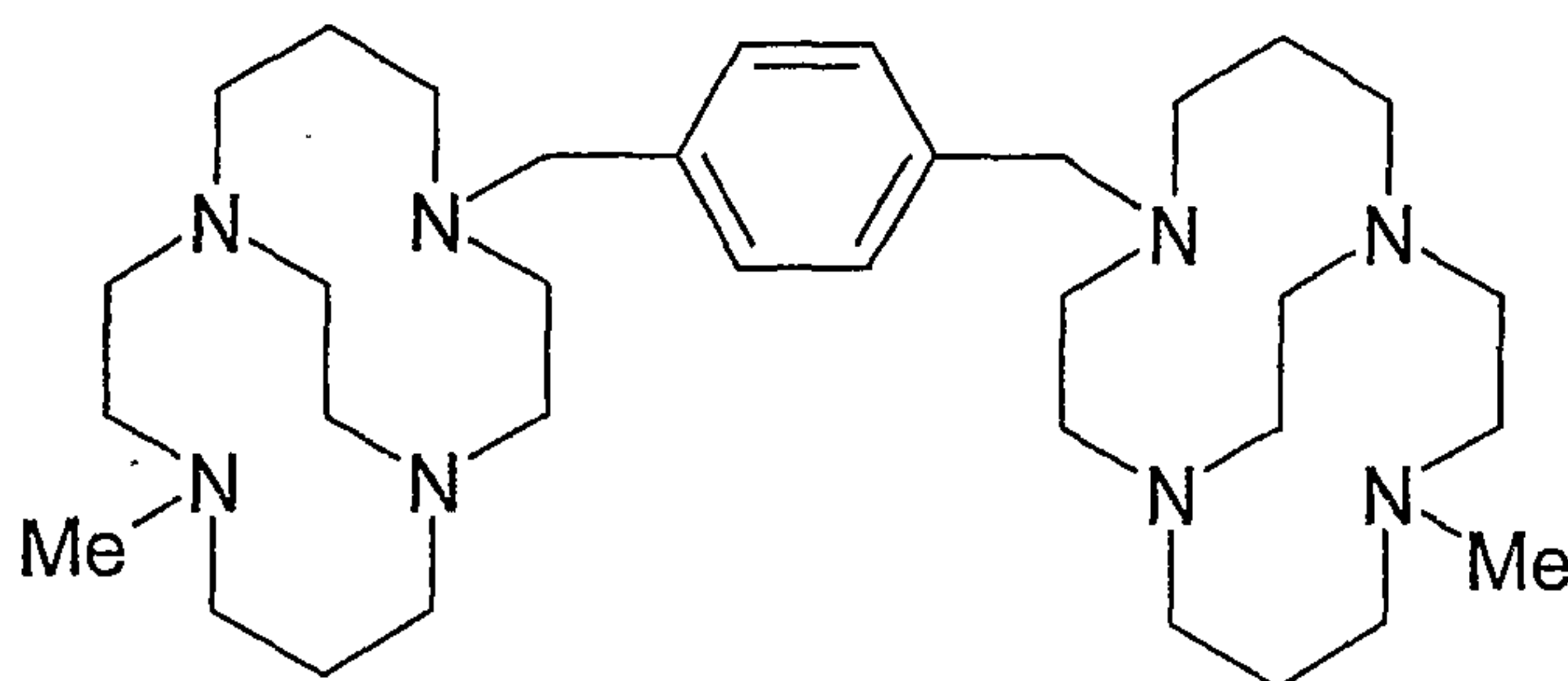
Example 2 Formula 17



para-Xylyl bis(1,5,8,12-tetraazabicyclo[10.2.2]hexadecane) (0.47g/0.85mmol) was dissolved in DCM (100ml). K_2CO_3 (0.22g/1.62mmol) and methyl iodide (0.23g/0.10ml/1.62mmol) were added and the mixture was stirred at room temperature under nitrogen overnight. The reaction mixture was filtered to remove K_2CO_3 and the filtrate was concentrated in vacuo to yield an orange solid. Mass and NMR (^1H and ^{13}C) spectra were consistent with the expected product.

Yield = 0.47g (96%).

Example 3 Formula 21



Synthesis of para-xylyl bis(1-methyl-1,5,8,12-tetraazabicyclo[6.6.2]hexadecane)

The bisaminal precursor, 0.41g (5.82 mmol) was added to 15ml of dry DMF and methyl iodide, 0.725ml (12.0 mmol) then the solution was stirred for 14 days. The solvent was removed on the rotary evaporator giving a white solid, (0.38g).

The white solid, 0.38g (6.6 mmol) in minimum of dry ethanol was reduced with NaBH_4 , 2.49g (66 mmol). The NaBH_4 was added over a period of 1 hour.

The reaction was left to stir for a further 14 days at room temperature. Excess NaBH_4 was decomposed with slow addition of 3M HCl (10ml) and the solvent removed. The resulting white solid was dissolved in water (30ml), the pH was adjusted to 14 using KOH pellets, and the basic solution was extracted with benzene (6x50ml).

The combined extracts were combined and the solvent removed, giving a yellow oil. Mass and NMR (^1H and ^{13}C) spectra were consistent with the expected product.

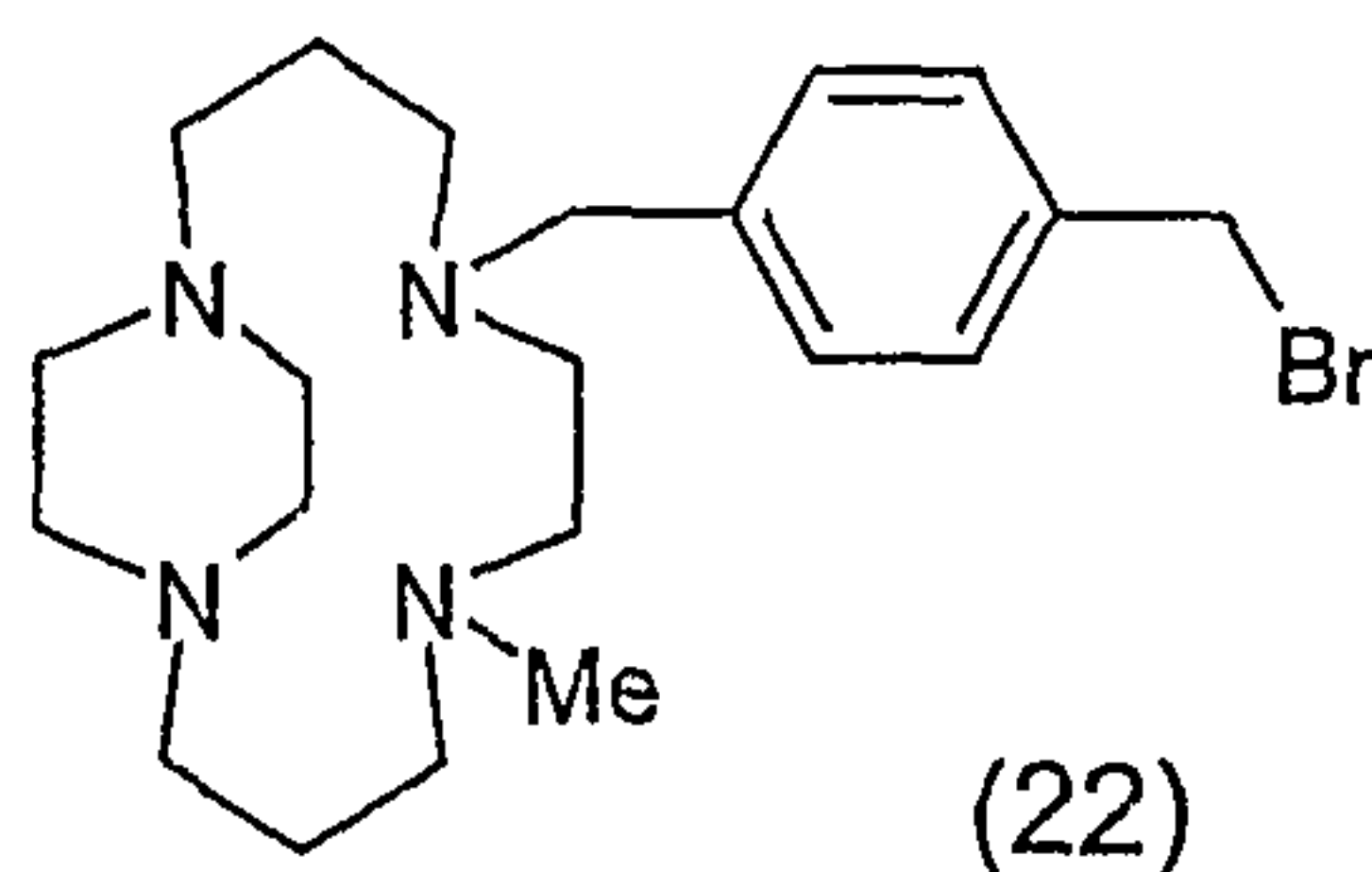
10 Example 4 Metal complexes Formulae 24 and 25

Ligand (112.4mg/0.2mmol) was dissolved in methanol (20ml) and copper chloride (0.07g/0.4mmol) was added. The blue solution was refluxed for 2 hours and stirred at room temperature for 48 hours. The solution was concentrated in vacuo and redissolved in a minimum amount methanol for recrystallisation.

15 Ligand (133.6mg/0.23mmol) was dissolved in methanol (20ml) and zinc acetate (0.10g/0.46mmol) was added. The orange solution was refluxed for 2 hours and stirred at room temperature for 48 hours. The solution was concentrated in vacuo and redissolved in a minimum amount methanol in an attempt to grow crystals.

Mass spectra and elemental analysis were consistent with the expected products.

Example 5 Unsymmetric compounds



25 Glyoxal bridged cyclam (4.97g/22.39mmol) was dissolved in dry THF (50ml) and 4-(methylbromo)benzoic acid ester (5.14g/22.39mmol) was added. The resulting yellow solution was stirred at room temperature under nitrogen for 4 days. The filtrate was decanted off and washed with further THF (4 x 20ml). The remaining precipitate was concentrated in vacuo to yield a white powder as product. Mass and NMR (^1H and ^{13}C) spectra were consistent with the expected product.

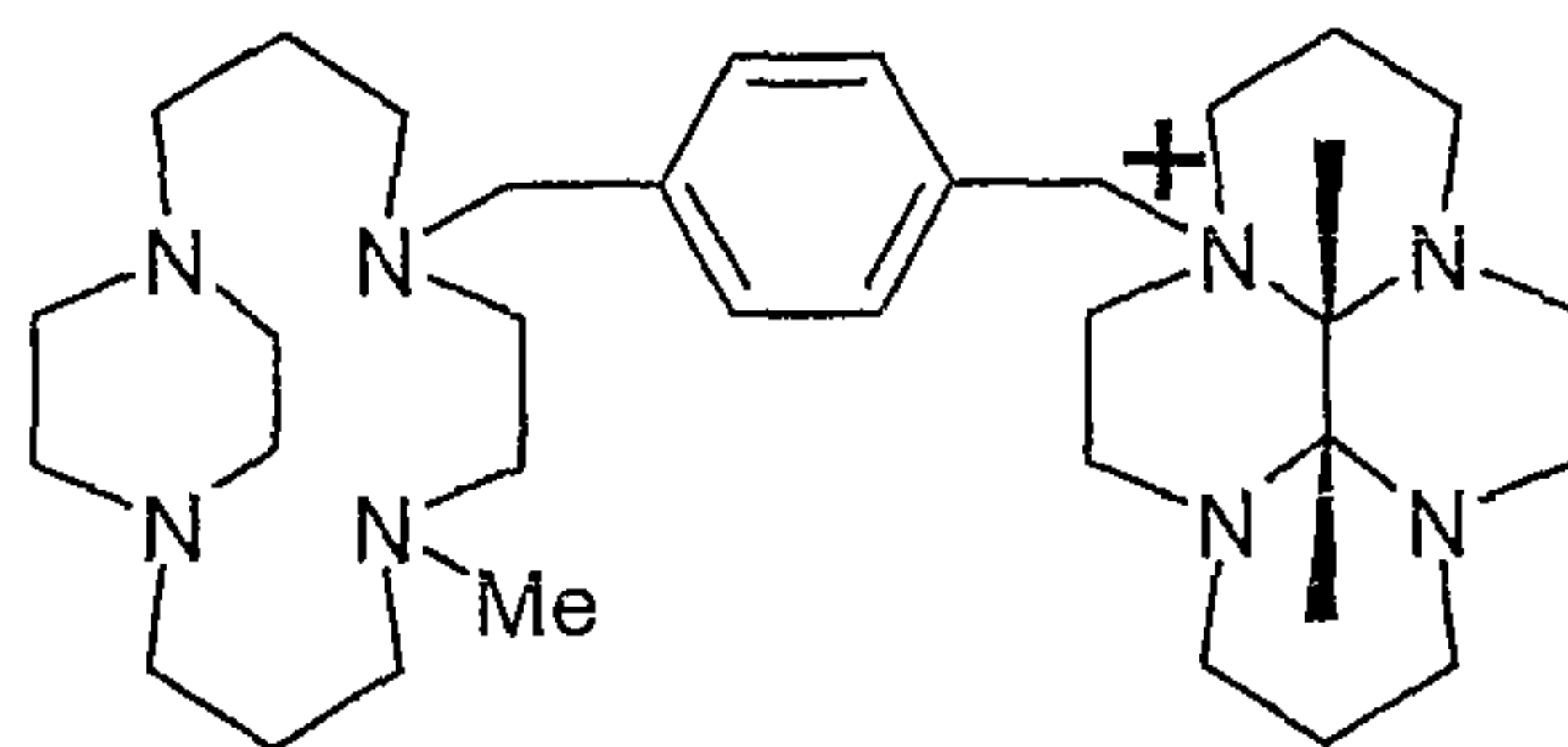
One-armed macrocycle (4.93g/10.91mmol) was dissolved in methanol (250ml) and cooled to 0°C in an ice bath. NaBH₄ (3.80g/100mmol) was added slowly over 30 minutes. The solution was stirred at room temperature for 1 hour and concentrated in vacuo to a white solid as the product. Mass and NMR (¹H and ¹³C) spectra were consistent with the expected product.

This compound (2.37g/6.35mmol) was dissolved in CH₃CN (50ml) and K₂CO₃ (0.87g/6.35mmol) was added. Methyl iodide (0.90g/0.40ml/6.35mmol) was added and the mixture was stirred at room temperature under nitrogen overnight. The resulting mixture was filtered to remove K₂CO₃ and concentrated in vacuo to yield an orange powder as product.

The one-armed ester (2.43g/6.35mmol) was dissolved in dry THF (100ml) and BH₃.SMe₂ (2.06ml/20.64mmol) was added. The mixture was refluxed under nitrogen for 6 days. The reaction was monitored by TLC to determine when it was complete. The reaction mixture was cooled in an ice bath and methanol was added until all the BH₃.SMe₂ had been consumed. The solution was concentrated in vacuo to yield a white solid. This was dissolved in brine (150ml) and extracted with dichloromethane (2 x 150ml, 1 x 300ml). The combined extracts were collected, dried (MgSO₄) and concentrated in vacuo to yield a white solid. This was purified by column chromatography (95% DCM, 5% methanol and 3 drops of Et₃N).

Yield = 0.93g (40%) (% yield for steps 4 and 5 together is 37%). ¹H NMR (CD₂Cl₂) δ 1.62-1.64 (m, 1H), 2.07 (s, 1H, CH₂OH), 2.40-2.79 (m, 6H), 3.28 (s, 5H), 3.61-3.66 (m, 2H), 3.88-3.89 (m, 3H), 4.60-4.70 (m, CH₂OH), 7.34-7.36 (m, 2H, CH_{aromatic}, J = 8.16Hz), 8.00-8.05 (m, 2H, CH_{aromatic} J = 8.16Hz). ¹³C NMR (CD₂Cl₂) δ 1.10, 67.18, 67.19, 77.40, 77.72, 78.04, 129.67 (C_{aromatic}), 129.96 (CH_{aromatic}), 130.00 (CH_{aromatic}).

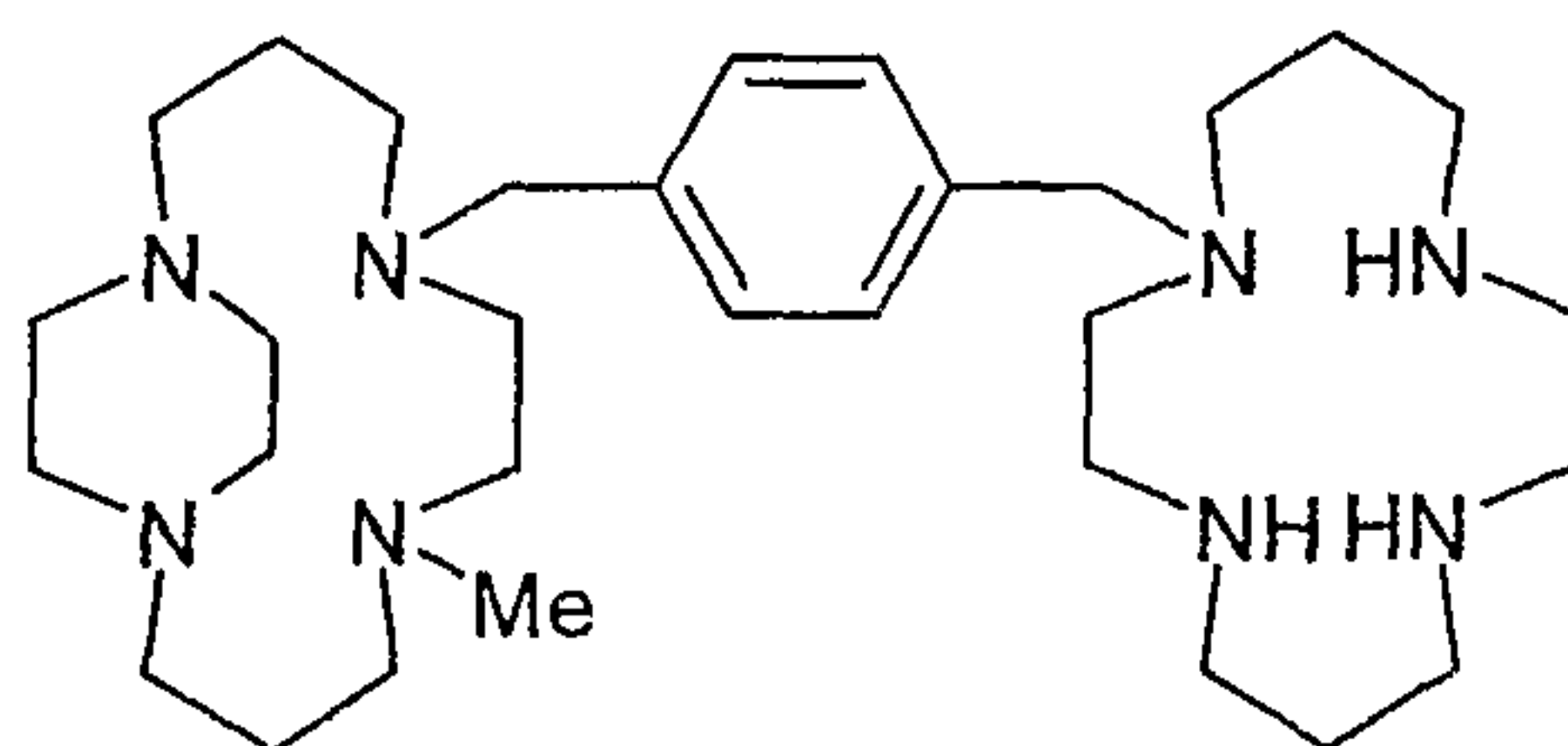
One-armed macrocycle alcohol (0.79g/2.20mmol) was added to PBr₃ (0.19ml/2.20mmol) and a catalytic amount of pyridine (~0.02ml) was added. The mixture was stirred at room temperature overnight (an orange colour was observed) and at 100°C for 1 hour. Water (20ml) was added slowly and made up to pH 14 with KOH pellets. The aqueous layer was extracted with dichloromethane (3 x 30ml). The combined extracts were collected, dried (MgSO₄) and concentrated in vacuo to yield an orange solid as product. Yield = 0.64g (69%) ¹H NMR (CD₂Cl₂, 400 MHz) δ: 7.19 (d, 8.2 Hz), 7.88 (d, 8.2 Hz), 3.17-3.00 (m, 2H), 2.97-2.83 (m, 2H), 2.82-2.68 (m, 2H), 2.60-2.18 (m, 16H), 2.15-2.02 (m, 2H), 1.72-1.50 (m, 6H).



(23)

One-armed bromide (0.30g/0.71mmol) was dissolved in CH₃CN (100ml) and butanedine bridged cyclam (0.18g/0.71mmol) was added. The resulting dark orange solution was stirred at room temperature under nitrogen for 48 hours. The filtrate was
 5 decanted off and washed with further CH₃CN (4 x 20ml). The remaining precipitate was concentrated in vacuo to yield a white/orange solid as product.

Yield = 0.23g (48%)



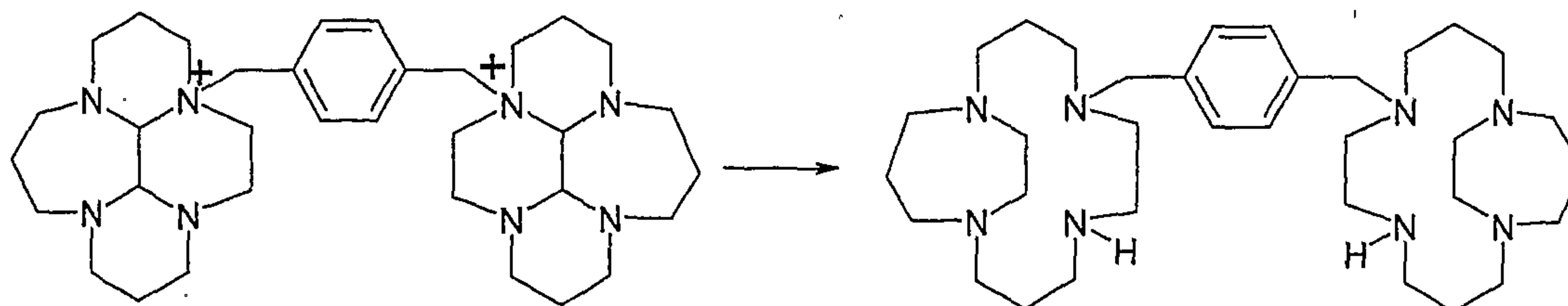
(20)

Bis-macrocycle with butanedione bridged pendent arm (11) (0.21g/0.31mmol)
 10 was dissolved in ethanol (50ml) and HCl (10% aq./50ml) was added. The orange solution was stirred at 60°C for 48 hours. The solution was concentrated in vacuo and water (50ml) was added. The pH was raised to 14 with KOH. The aqueous layer was extracted with DCM (3 x 100ml). The combined extracts were collected, dried (MgSO₄) and concentrated in vacuo to yield a yellow oil.

15

Example 6 1,5,9,13-Tetraazabicyclo[11.2.2]heptadecane derivatives

1,1'-[1,4-phenylenebis(methylene)]-bis(1,5,9,12 Tetraazabicyclo[3.6.9.11]heptadecane)



20

The bis-monoquaternary macrocycle salt [synthesised by a modified procedure from M. Le Baccon *et al*, New Journal of Chemistry, 2001, 25, 118.] (1.27 mmol, 0.933g) in dry ethanol (100ml) was cooled in an ice bath under an inert atmosphere. Sodium borohydride (12.2 mmol, 0.479g) was added slowly. The mixture was stirred
5 for 30 minutes and refluxed for 1 hour. The solvent was evaporated and water (60ml) added. The pH was then raised to 14 using KOH. Dichloromethane was used to extract the basic solution and the extracts were dried over sodium sulphate for 30 minutes. The solution was then filtered and the solvent removed from the filtrate to give a white solid. Mass and NMR (^1H and ^{13}C) spectra were consistent with the
10 expected product. Yield = 91%.

Formation of the HCl salt.

The bis(macrocycle) (0.3448mmol, 0.2g) was dissolved in ethanol (20ml) and HCl gas was bubbled through the solution for 5 minutes. A white powder precipitated
15 out and was filtered off to yield the HCl salt. Mass and NMR (^1H and ^{13}C) spectra were consistent with the expected product. Yield = 85%

Formation of a copper complex

Copper acetate (0.74 mmol, 0.158g) in methanol (30ml) was slowly added to
20 stirred solution of the bis(macrocycle) (0.34 mmol, 0.20g) in methanol (20ml) under an inert atmosphere. The copper solution changed colour from turquoise to blue on addition to the macrocycle solution, indicating that the copper was complexing with the macrocycle. The mixture was stirred and refluxed for 1 hour. After which it was dried on the schlenk line to yield a blue crystalline powder. The mass spectrum was
25 consistent with the expected product. Yield = 0.29g (90%)

Formation of the zinc complex

Zinc acetate (0.74 mmol, 0.159g) in methanol (30ml) was slowly added to
stirred solution of the bis(macrocycle) (0.34 mmol, 0.20g) in methanol (20ml) under an inert atmosphere. The mixture was stirred and refluxed for 1 hour after which the
30 product was dried on the schlenk line to yield the zinc complex. The mass spectrum was consistent with the expected product.

Example 7Fluorophore tagged CXCR4 binding molecules for receptor imaging*Formation of Rhodamine-bridged macrocycle conjugate*

- 5 The conjugate was formed following the typical procedure for combining an isothiocyanate with a free amine group to form a thiourea linkage [example methodology M. M. Hüber *et al.*, Bioconjugate Chemistry, 1998, 9, 242.]. Equimolar amounts of rhodamine B isothiocyanate was combined with the one armed macrocycle containing a 4-aminobenzyl moiety and triethylamine in methanol and
- 10 stirred at RT overnight. The product was purified by passing down a sephadex LH-20 column eluted with methanol. The NMR and mass spectral data were consistent with the desired conjugate.

Formation of copper complex

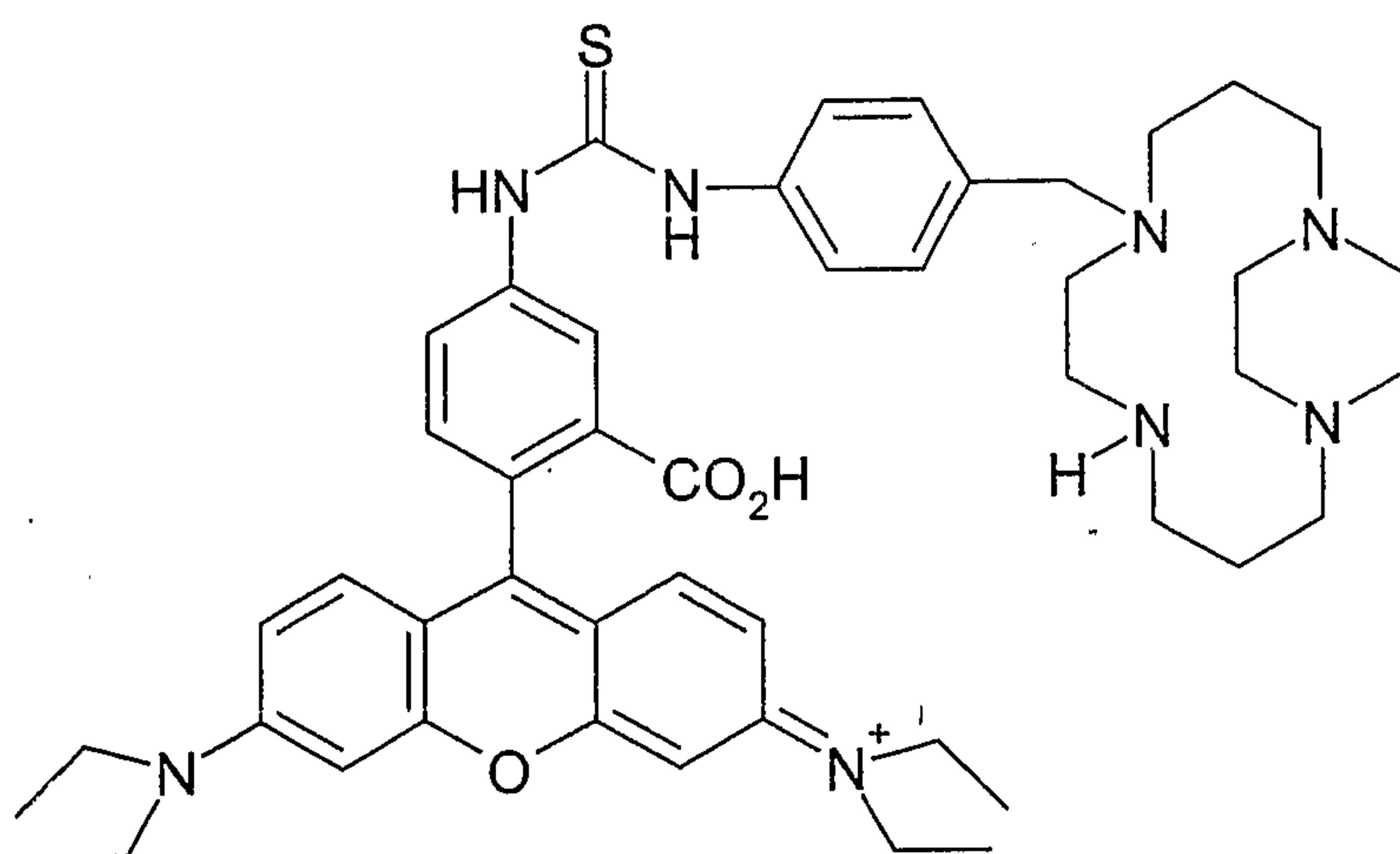
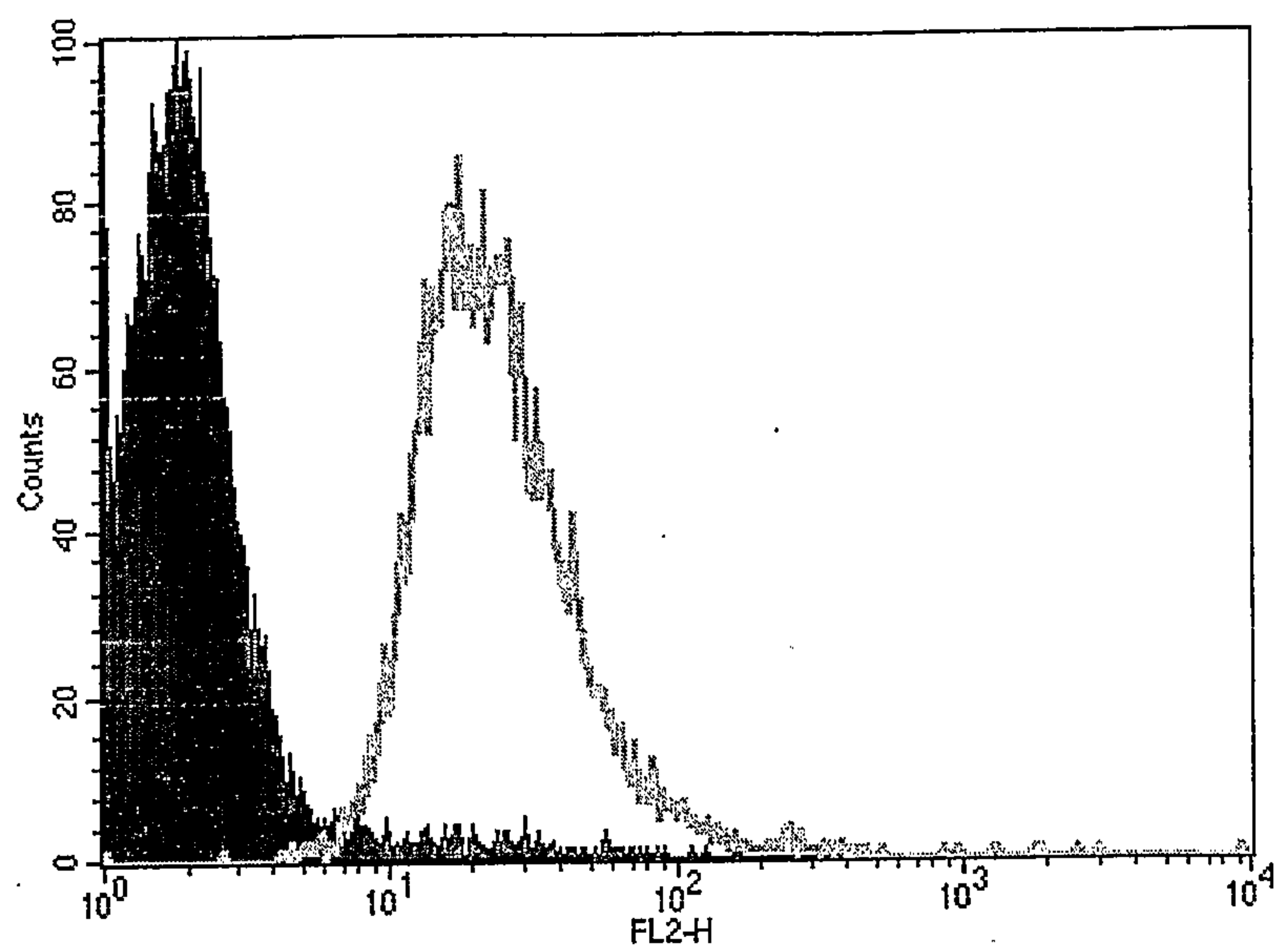
- 15 Equimolar amounts of copper perchlorate and rhodamine B-bridged macrocycle conjugate were combined in ethanol and heated to reflux for 2 h. On evaporation the complex was obtained as a purple solid that could be purified further by passing down a sephadex LH-20 plug in methanol.

20 *Binding of Rhodamine-bridged macrocycle conjugates and complexes*

Two monocyclam compounds were tagged with rhodamine and specific binding to the CXCR4 receptor was analysed by incubating with jurkat cells. Cell samples were analysed by flow cytometry as described in example 8, below.

25 *Rhodamine-bridged macrocycle conjugate activity*

This shows the structure of a novel rhodamine fluorophore conjugate molecules and the flow cell cytometry binding data showing binding of the chelator to the CXCR4 chemokine receptor.



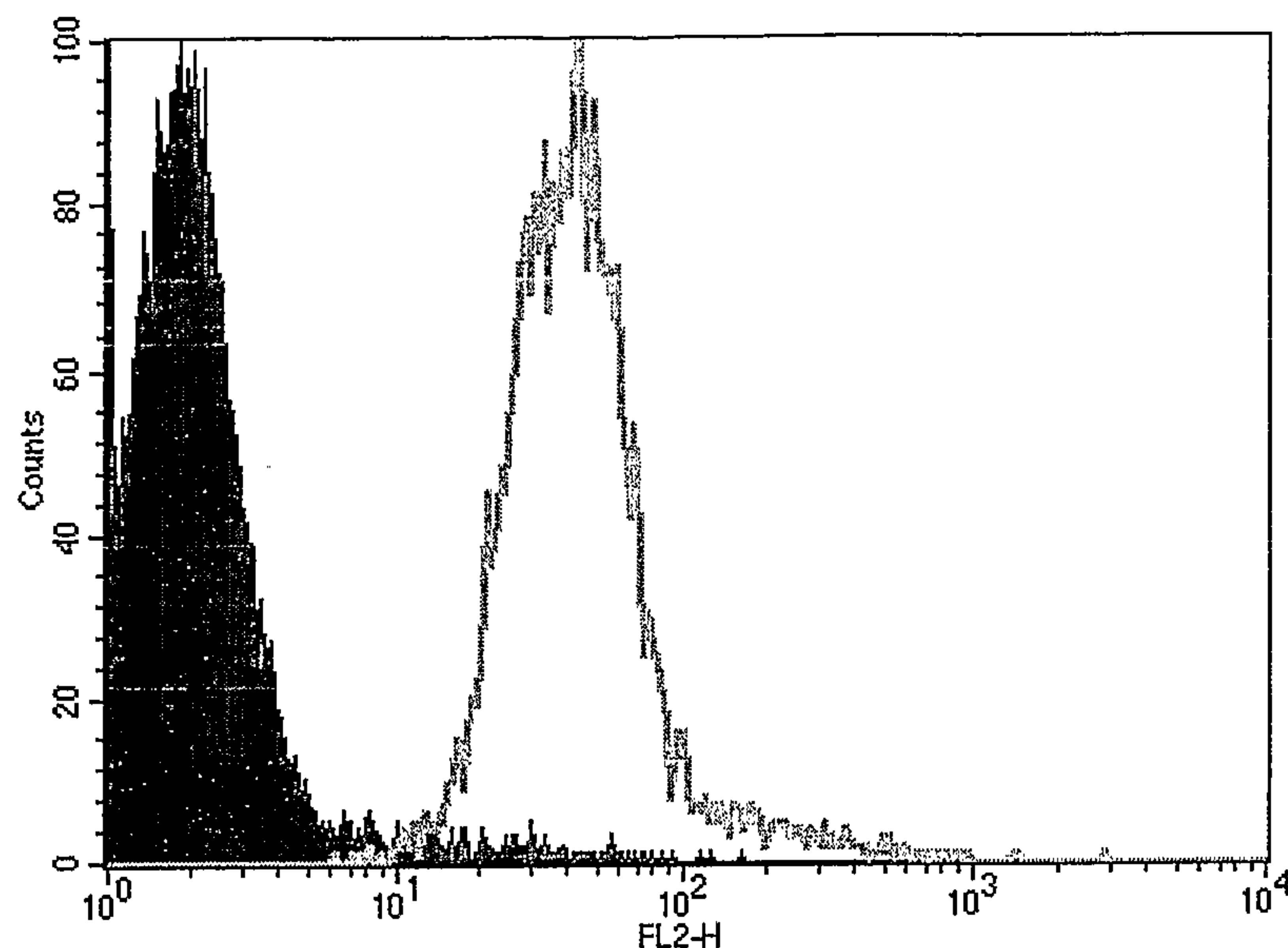
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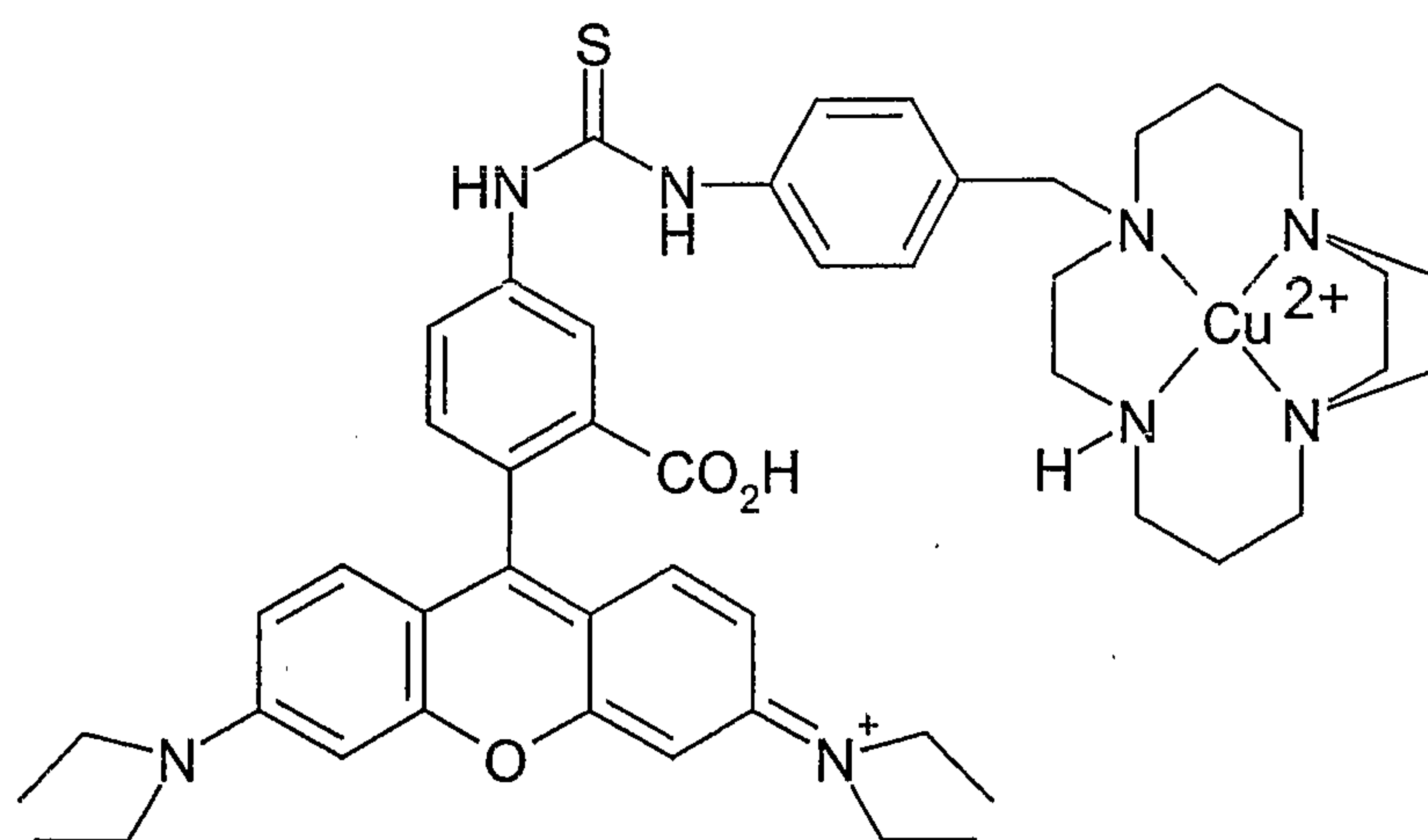
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Rhodamine-bridged macrocycle copper complex

This shows the structure of a novel rhodamine fluorophore conjugate molecule and the flow cell cytometry binding data showing binding of the copper complex to the CXCR4 chemokine



5 receptor.



Example 8

10 Competition with CXCR4 binding antibodies demonstrating CXCR4 receptor binding

Antibody stainings and flow cytometry

The antibodies used in this study were: unconjugated mouse anti-human CXCR4 mAbs clones 44708.111, 44716.111, PE-conjugated clone 44717.111 and
 15 12G5 (R&D Systems Europe, Abingdon, UK). Cells were preincubated with 20μM

of the compound for 30 mins. Thereafter cells were incubated with each of the mAbs for a further 60 mins. In the case of the direct binding assay, serial dilutions of the compound were added simultaneously with the antibody (44708.111). Cells were then incubated with a secondary fluorescein isothiocyanate-conjugated anti-mouse
 5 antibody (IgG-FITC)(Serotec, UK) for 30 mins. Cell samples were analysed by a FACScan flow cytometer (BD Biosciences Europe, Erembodegem, Belgium). Data were acquired and analysed with CellQuest software (Becton Dickinson) on an Apple Macintosh computer. Relative percentage inhibitions were calculated from the mean
 10 fluorescent intensity of the whole cell population and relative inhibitory concentration (IC) values were obtained by non linear regression analyses using the Graphpad prism software.

The following table shows antibody competition studies that demonstrate our molecules effectively bind to the CXCR4 chemokine receptor and compete with
 15 CXCR4 specific monoclonal antibodies.

Antagonist	IC50 (μ M)	IC90 (μ M)
25	0.178	0.853
Zn(21)	0.254	1.349
24	0.260	1.516
Cu(16)	0.331	2.845
16	17.025	>17.025

Anti-viral inhibitory action and cytotoxicities of examples of the drugs

20 *Antiviral Assay and Cytotoxicity Assay*

Anti-HIV activity and cytotoxicity measurements in MT-4 cells were based on viability of cells that had been infected or not infected with HIV-1 (IIIB) and HIV-2 (ROD) x4 strainsexposed to various concentrations of the test compound [following the procedure of S. Harada *et al.*, Science, 1985, 229, 563]. The MT-4 cells were
 25 allowed to proliferate for 5 d after which time the number of viable cells was

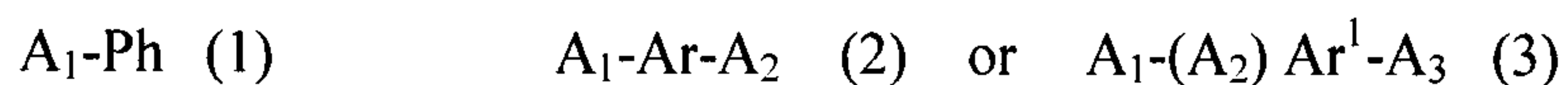
quantified by a tetrazolium-based colorimetric method [described by R. Pauwels *et al.*, Journal of Virology Methods, 1988, 20, 309.]

5 This data demonstrates the in-vitro anti-HIV activity of some of our compounds showing inhibition of infection by the HIV virus in MT-4 cells.

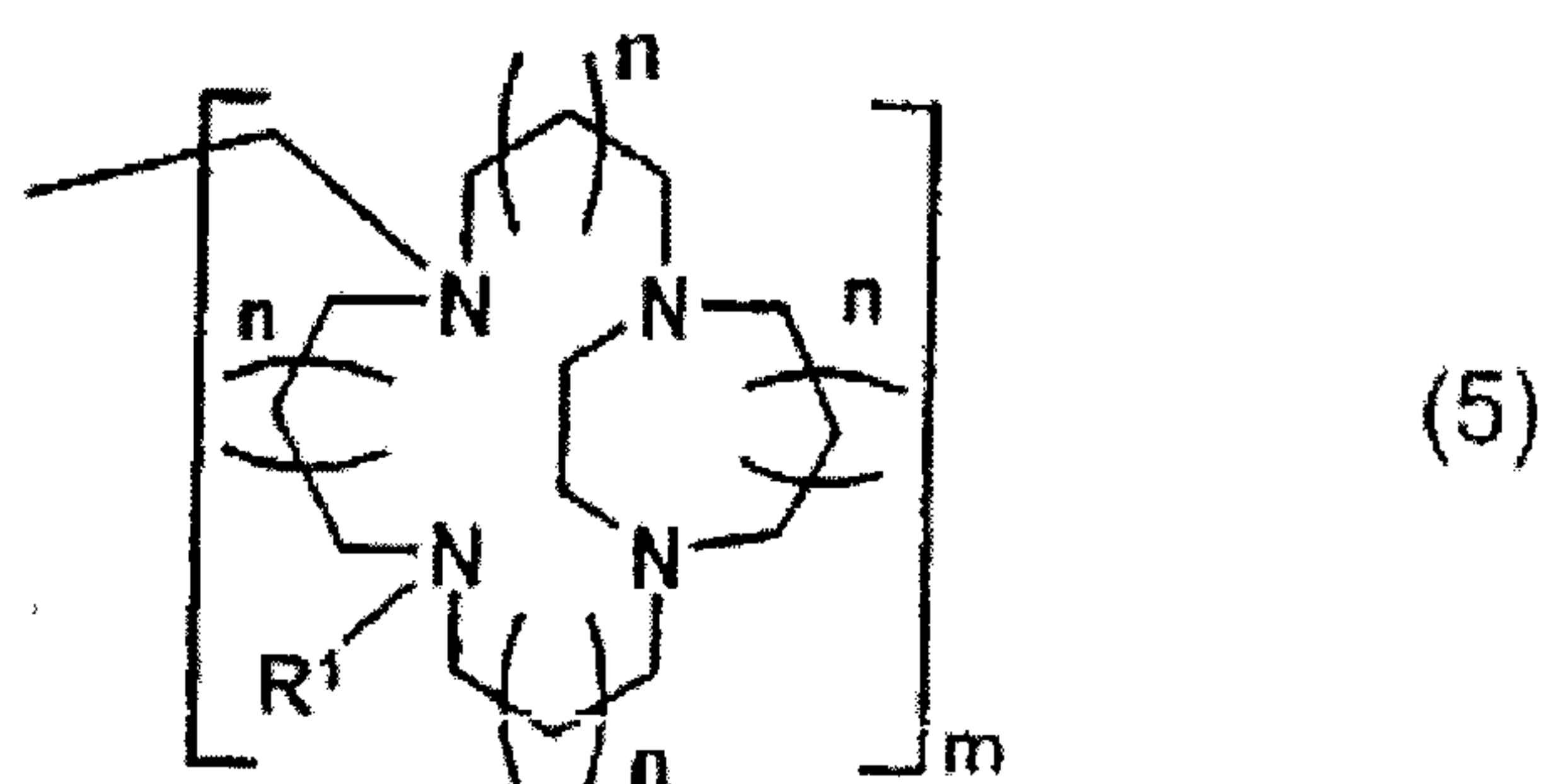
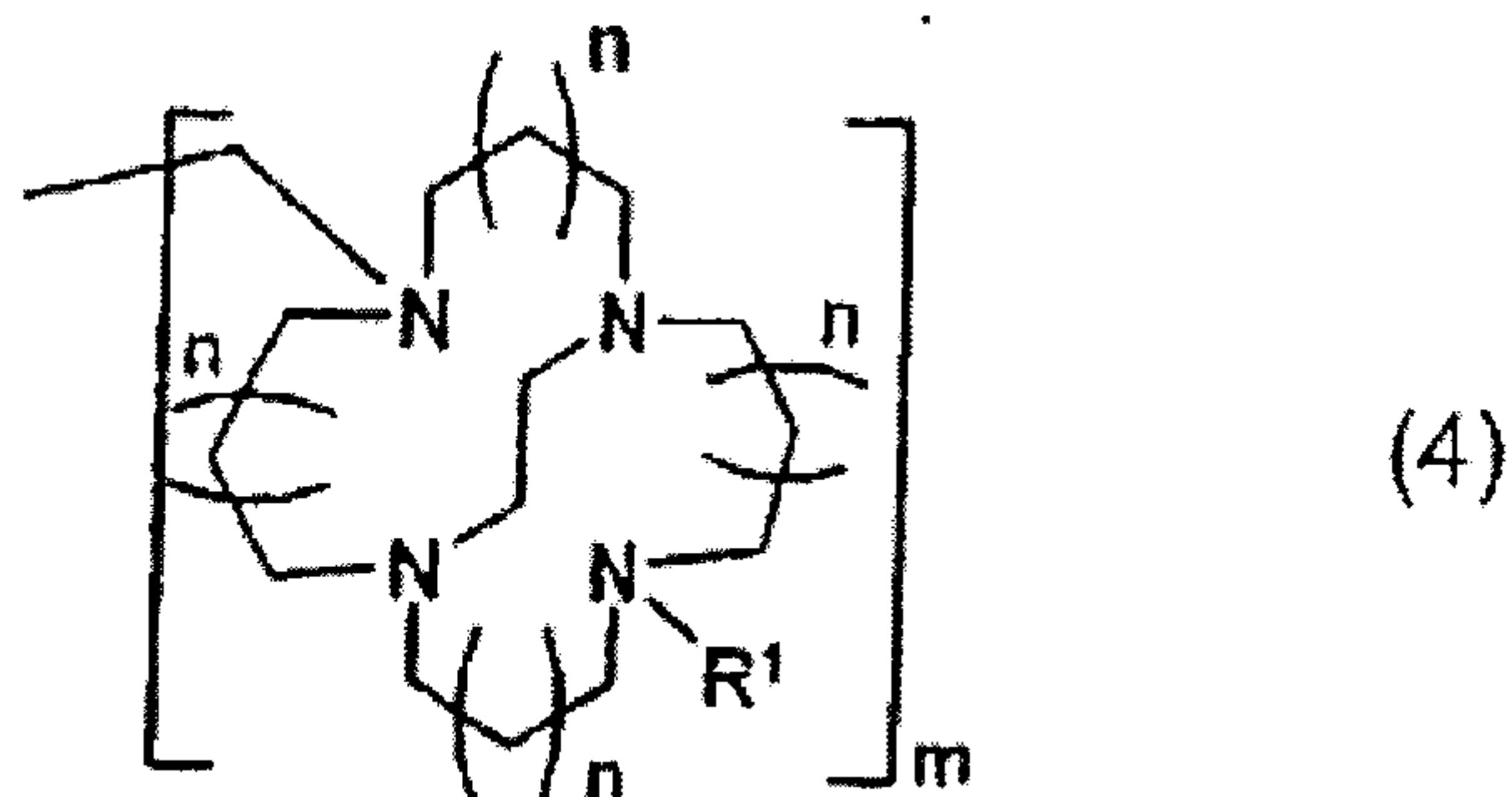
Compound	HIV strain	av IC50(μ M)	av CC50(μ M)
16	HIV-1	6.98	>225
	HIV-2	23.2	>225
Cu(16)	HIV-1	0.028	>150
	HIV-2	0.060	>150
24	HIV-1	0.0025	60.56
	HIV-2	0.0040	60.56

CLAIMS:

1. A compound of formula 1, 2 or 3,



wherein A_1 , A_2 and A_3 are selected from formulae 4 and 5



or a salt thereof;

wherein Ar is selected from the group consisting of 1,2-phenyl, 1,3-phenyl and 1,4-phenyl optionally substituted with one or more methyl groups;

each n is independently an integer from 0 to 1;

m is 1;

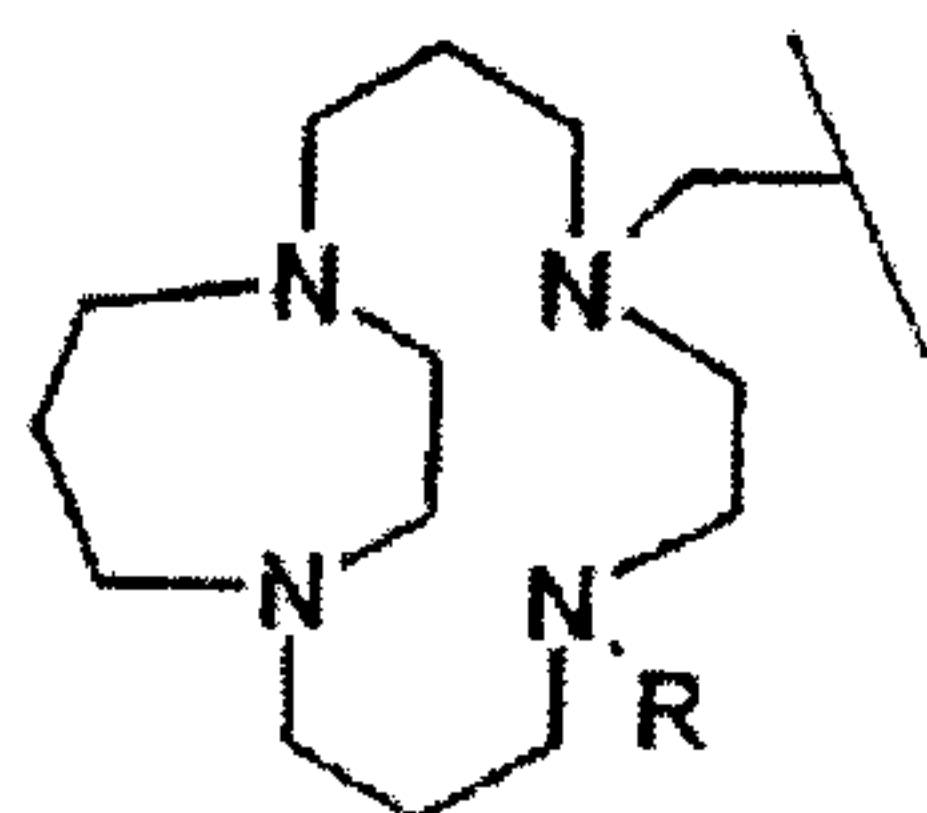
Ph is phenyl optionally functionalised with groups R^4 , wherein R^4 is selected from the group consisting of methyl, other C_1 to C_4 alkyl and fluorophores;

Ar^1 is trisubstituted phenyl optionally substituted with one or more phenyl groups; and

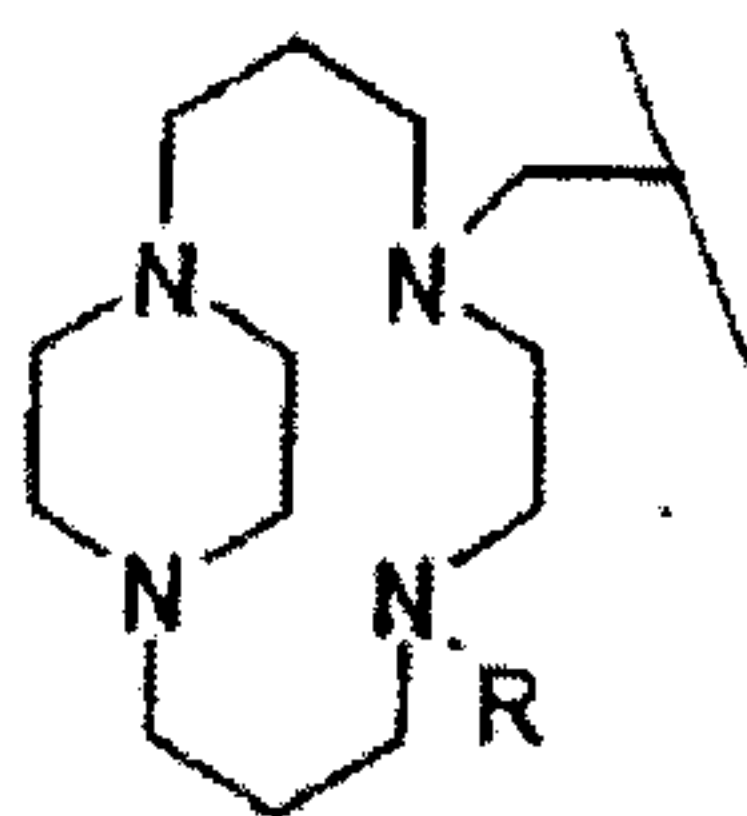
R^1 is selected from the group consisting of H, Me and linear chain C_2 - C_4 alkyl;

with the proviso that when Ph is unsubstituted phenyl in formula 1 and A_1 is formula 4, R_1 is not Me.

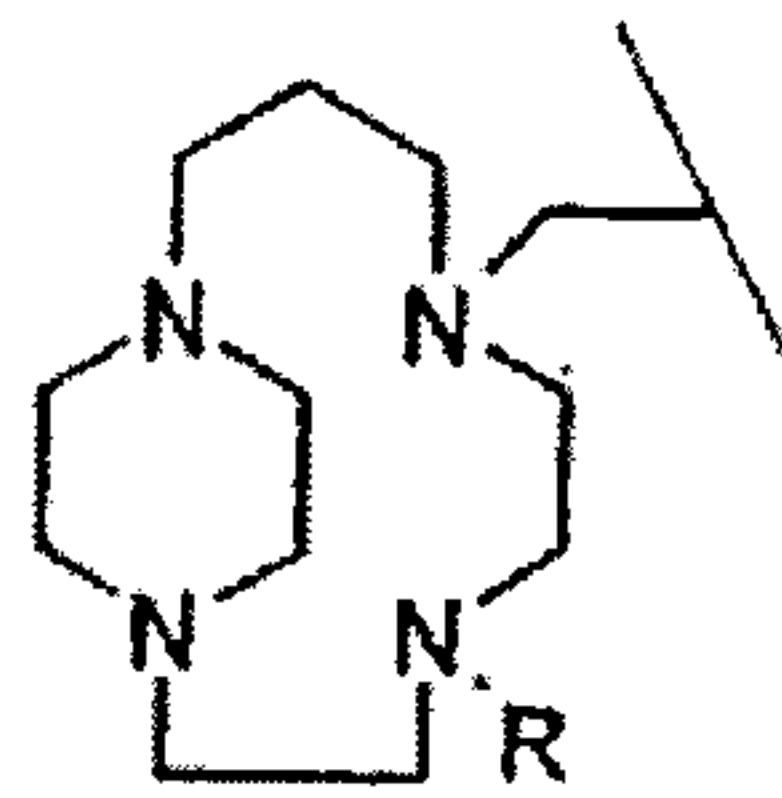
2. A compound as claimed in claim 1 wherein A_1 , A_2 and A_3 are independently selected from a group consisting of the formulae 7 to 15,



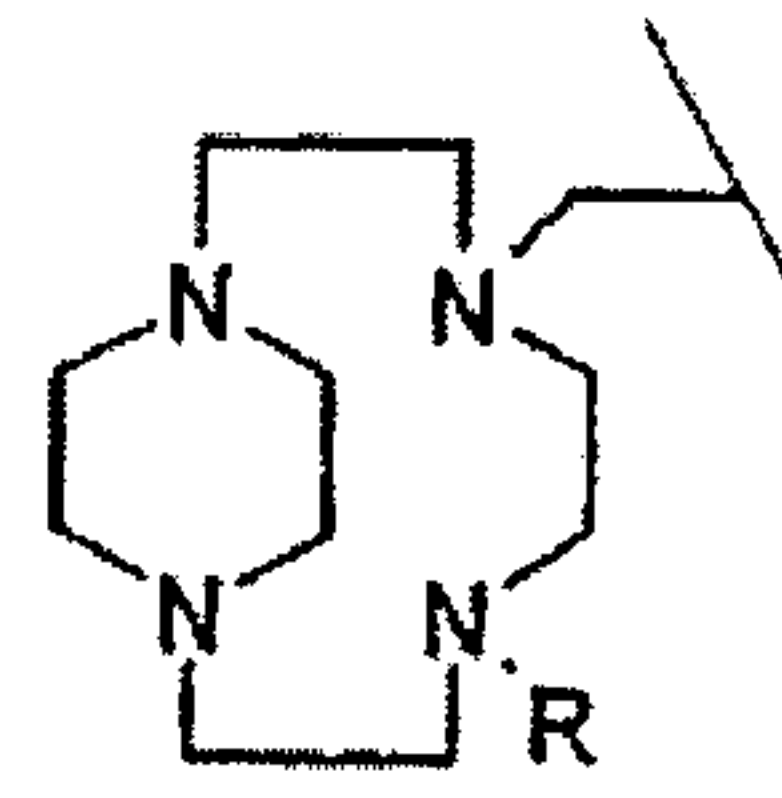
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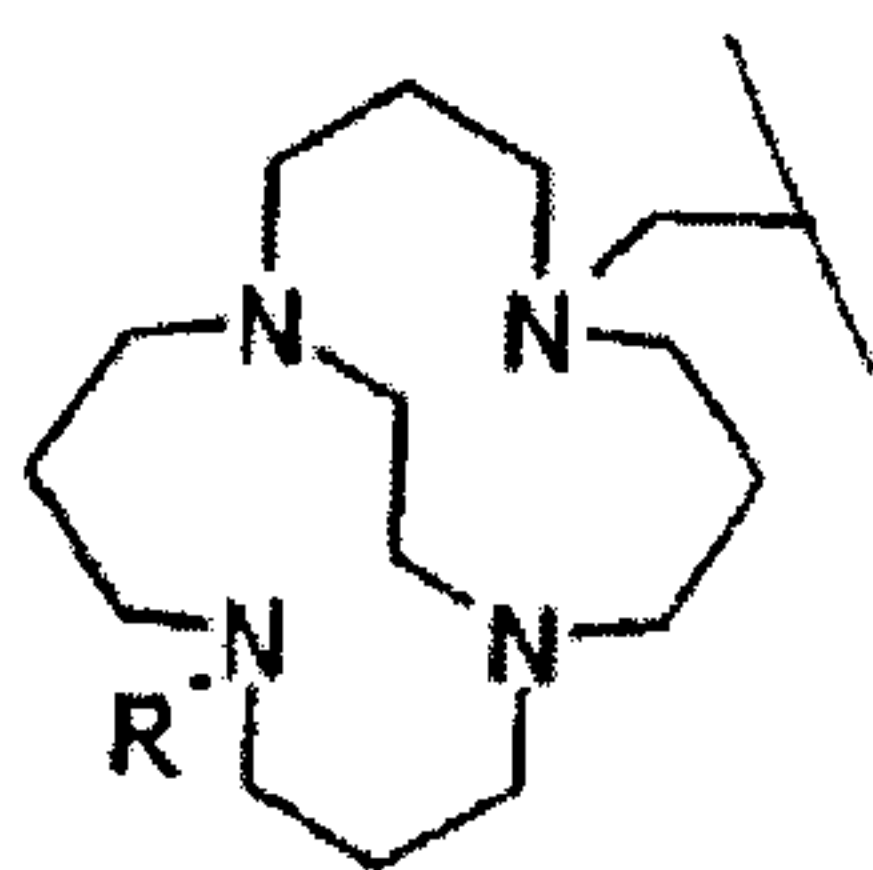
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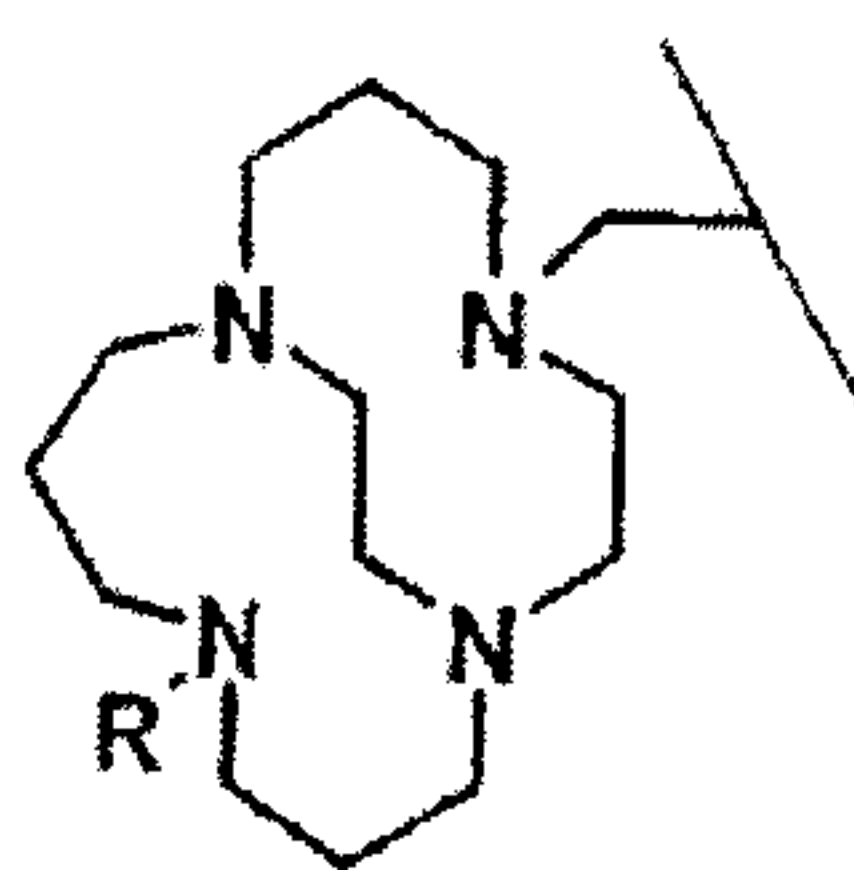
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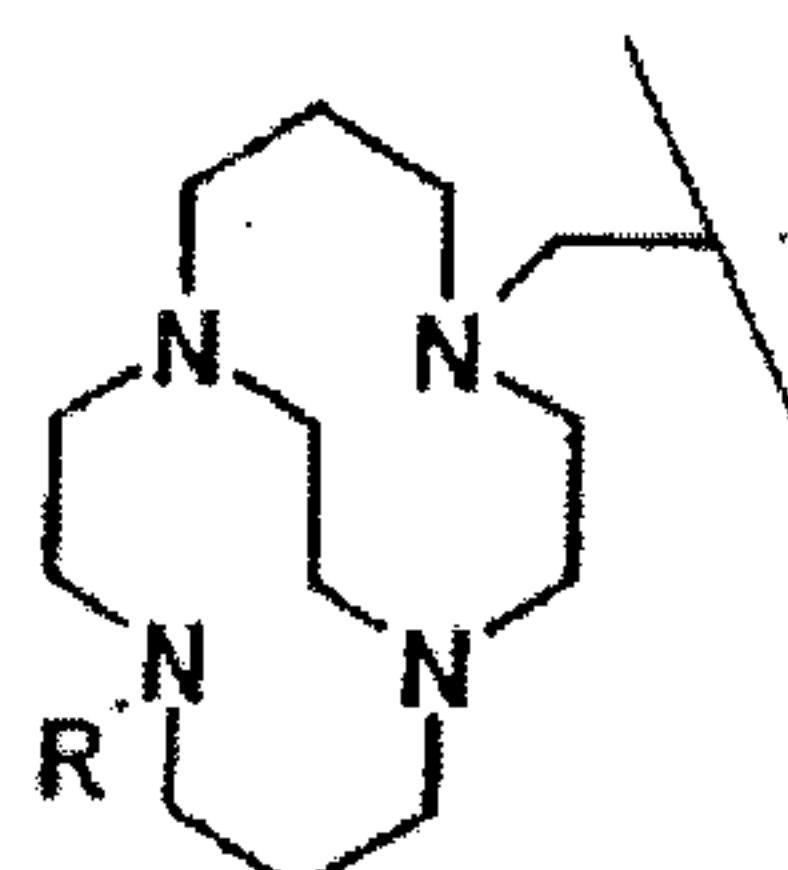
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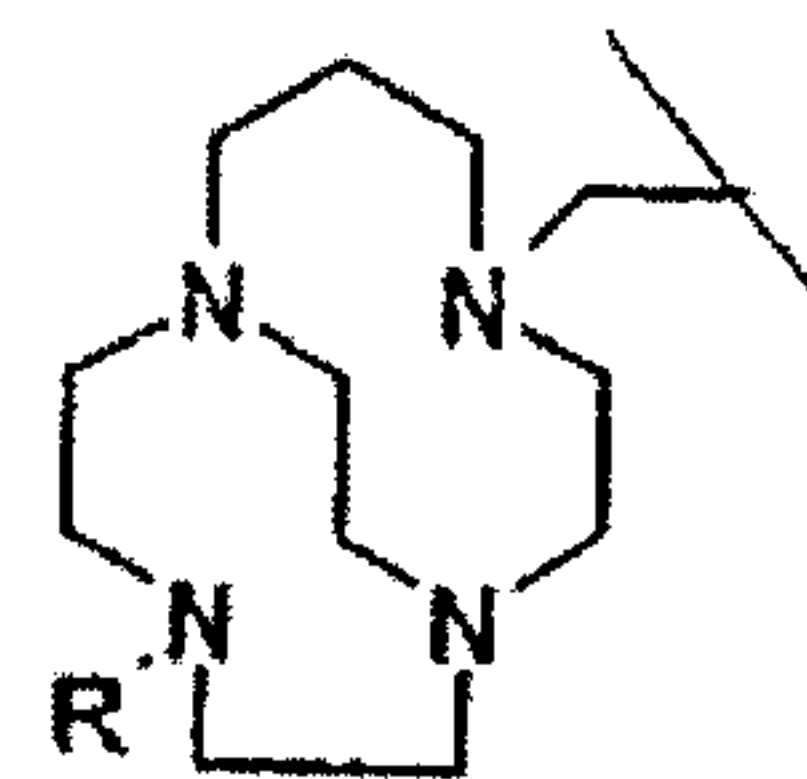
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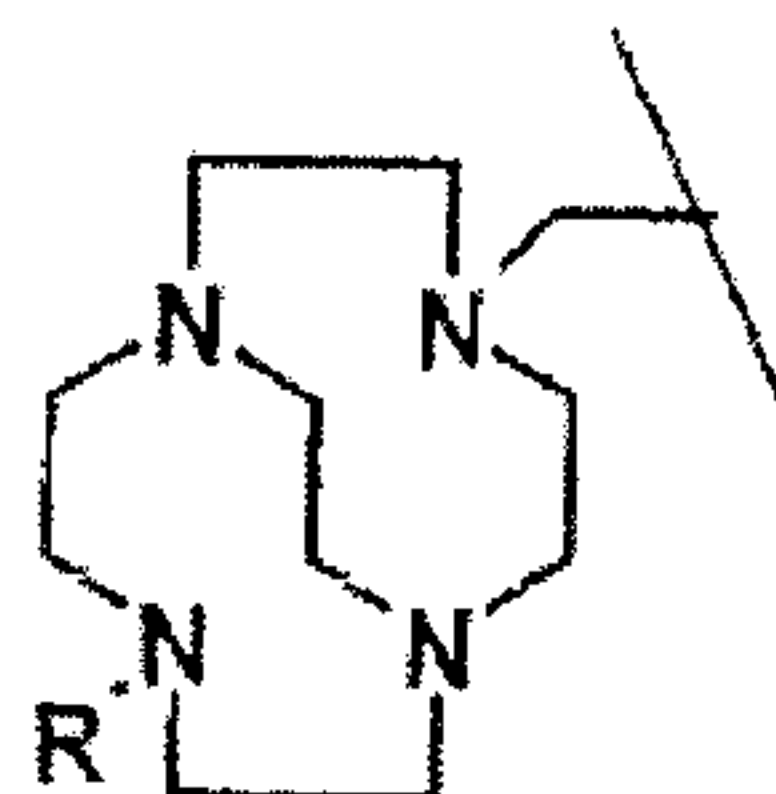
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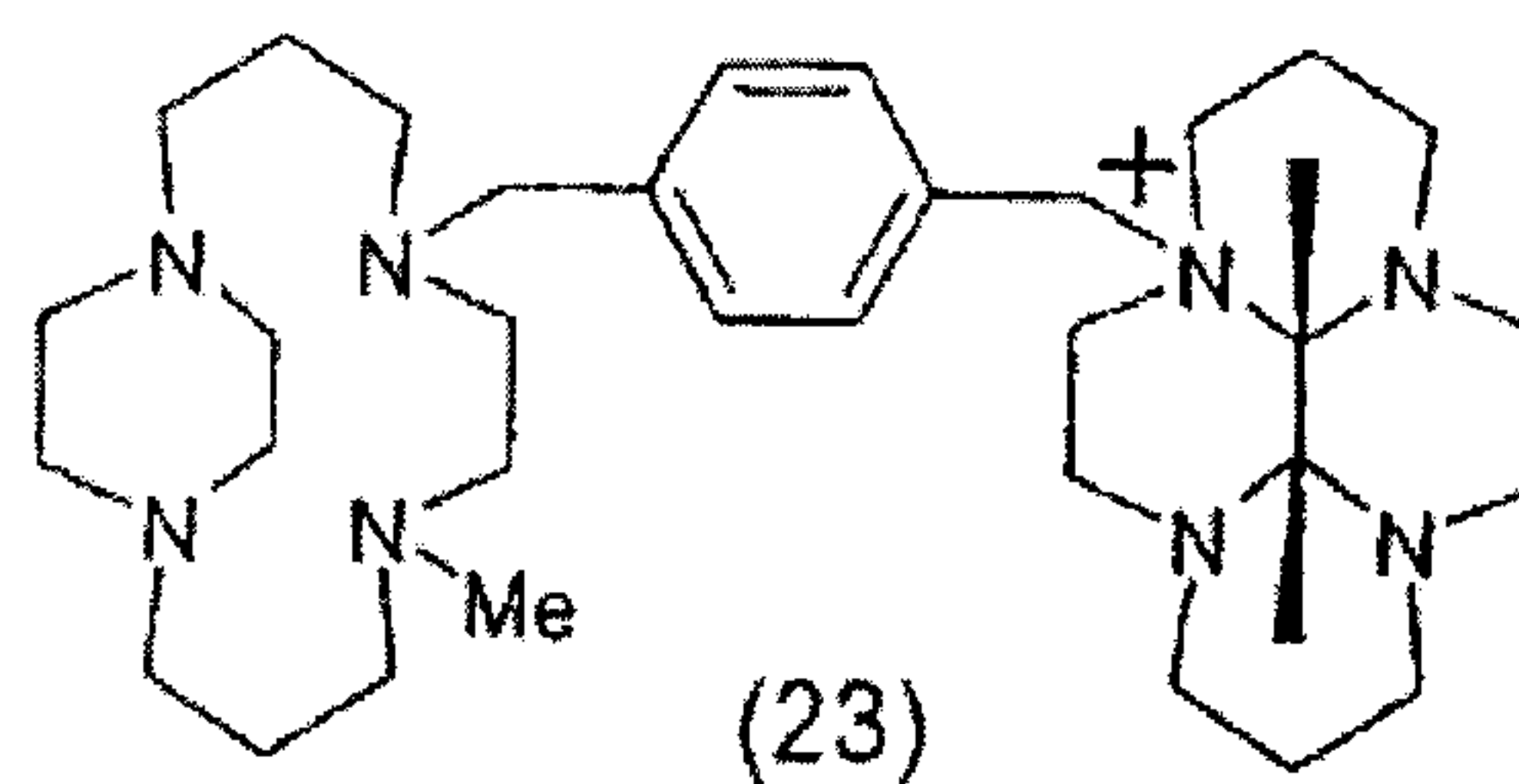
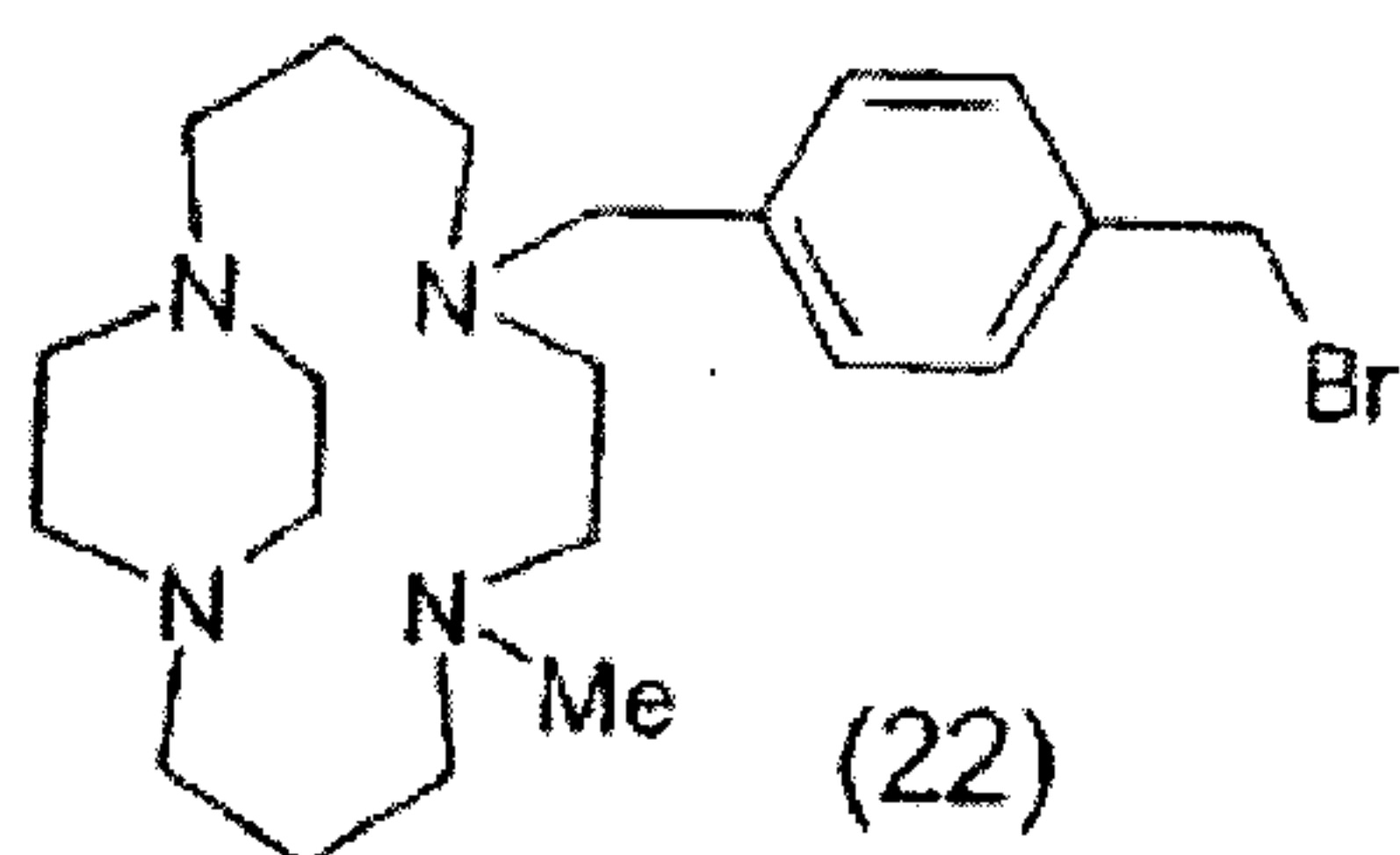
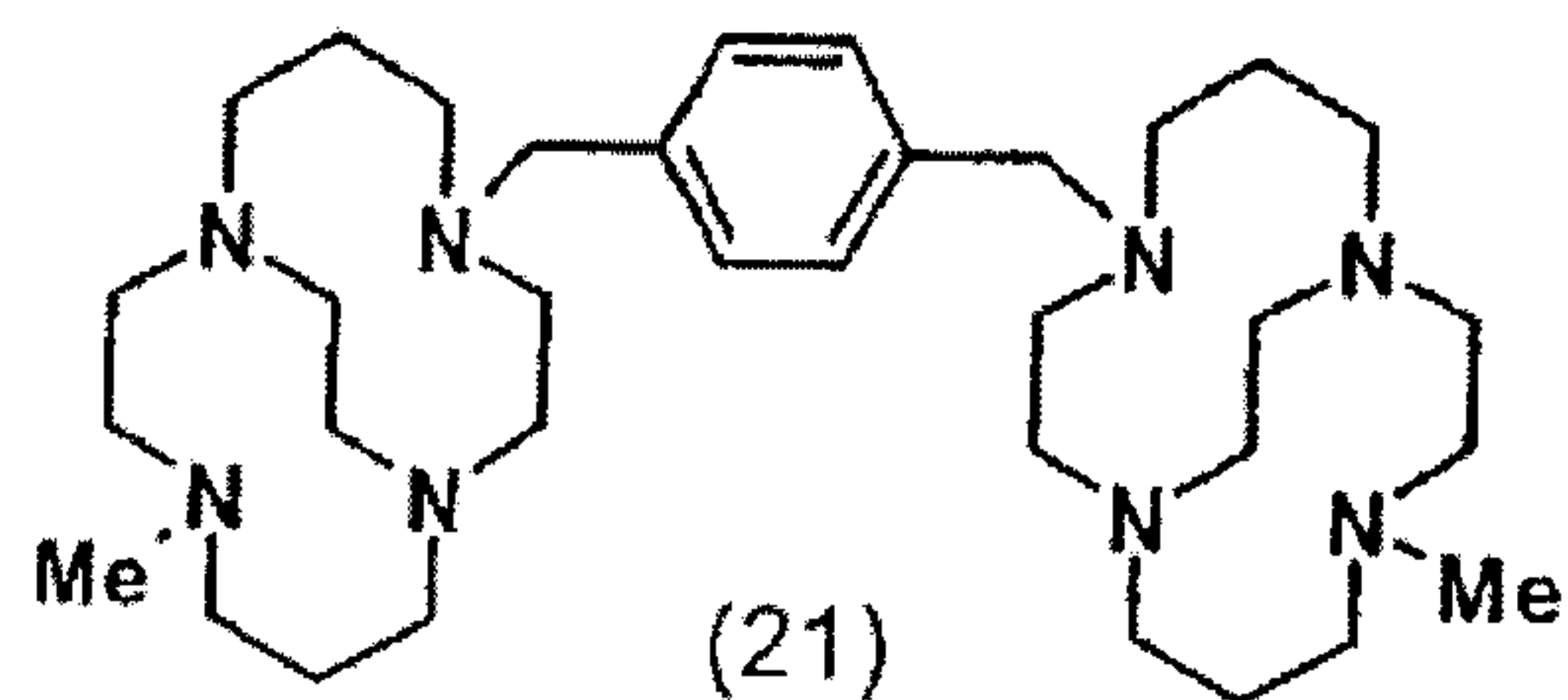
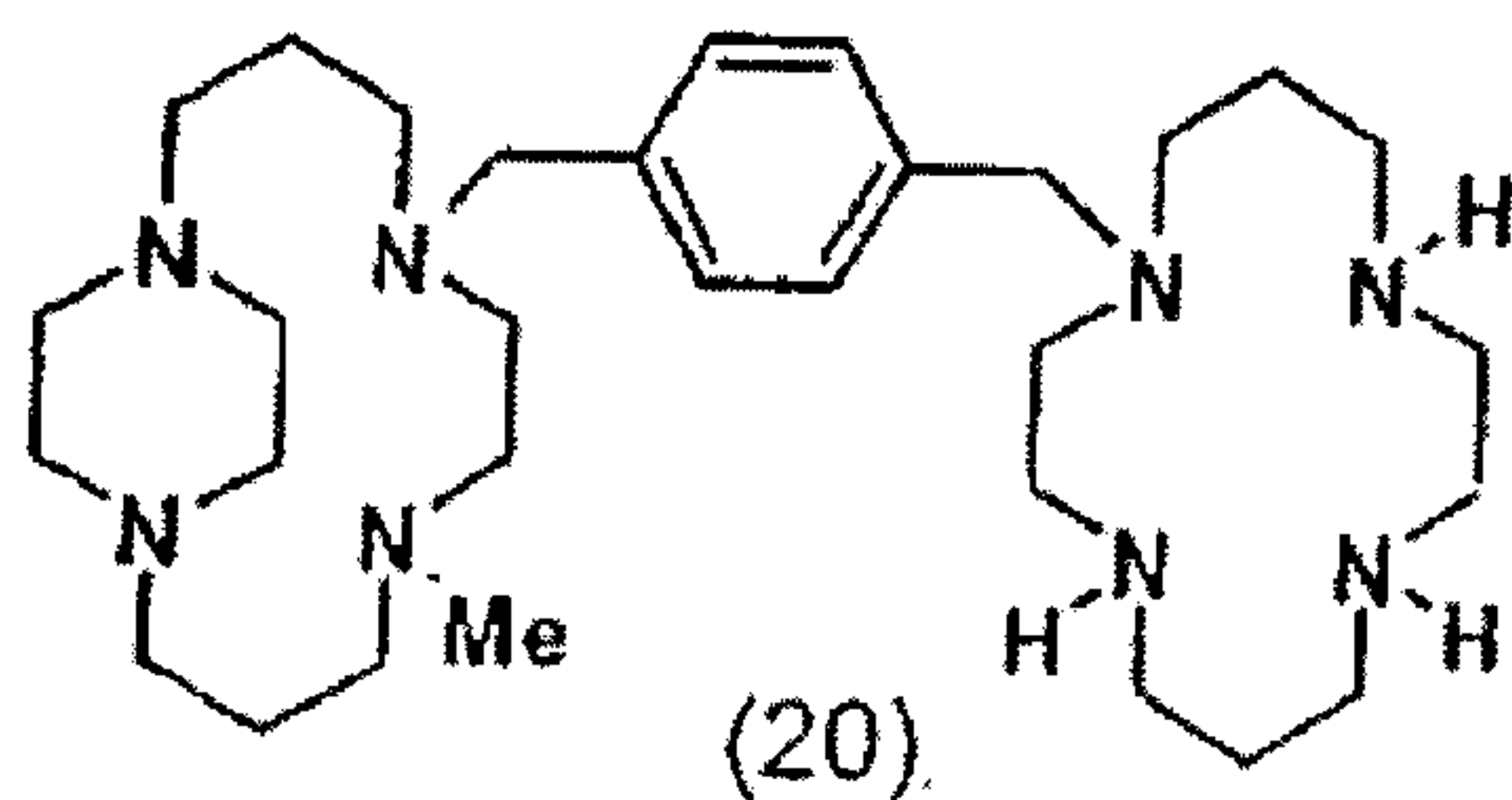
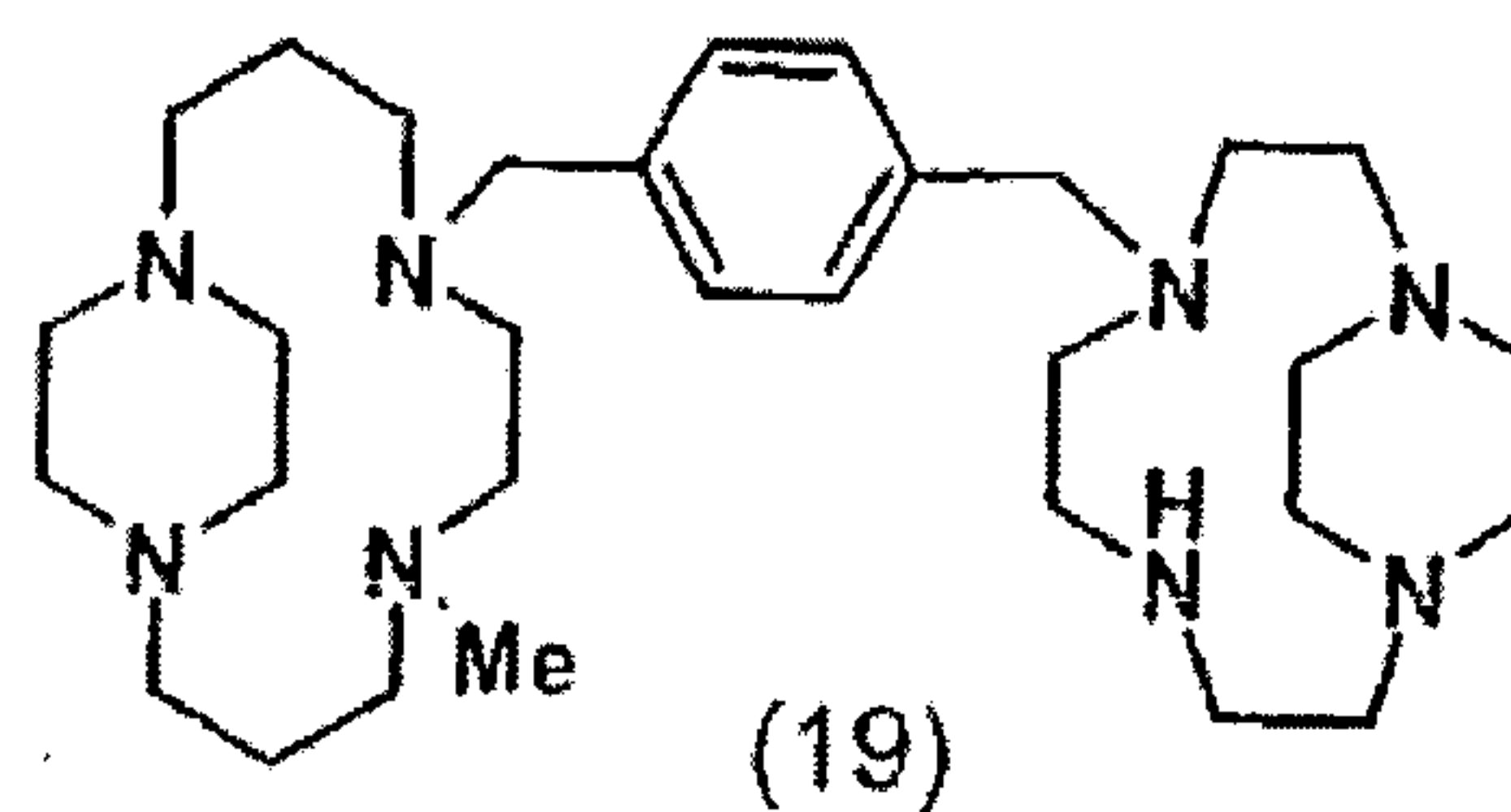
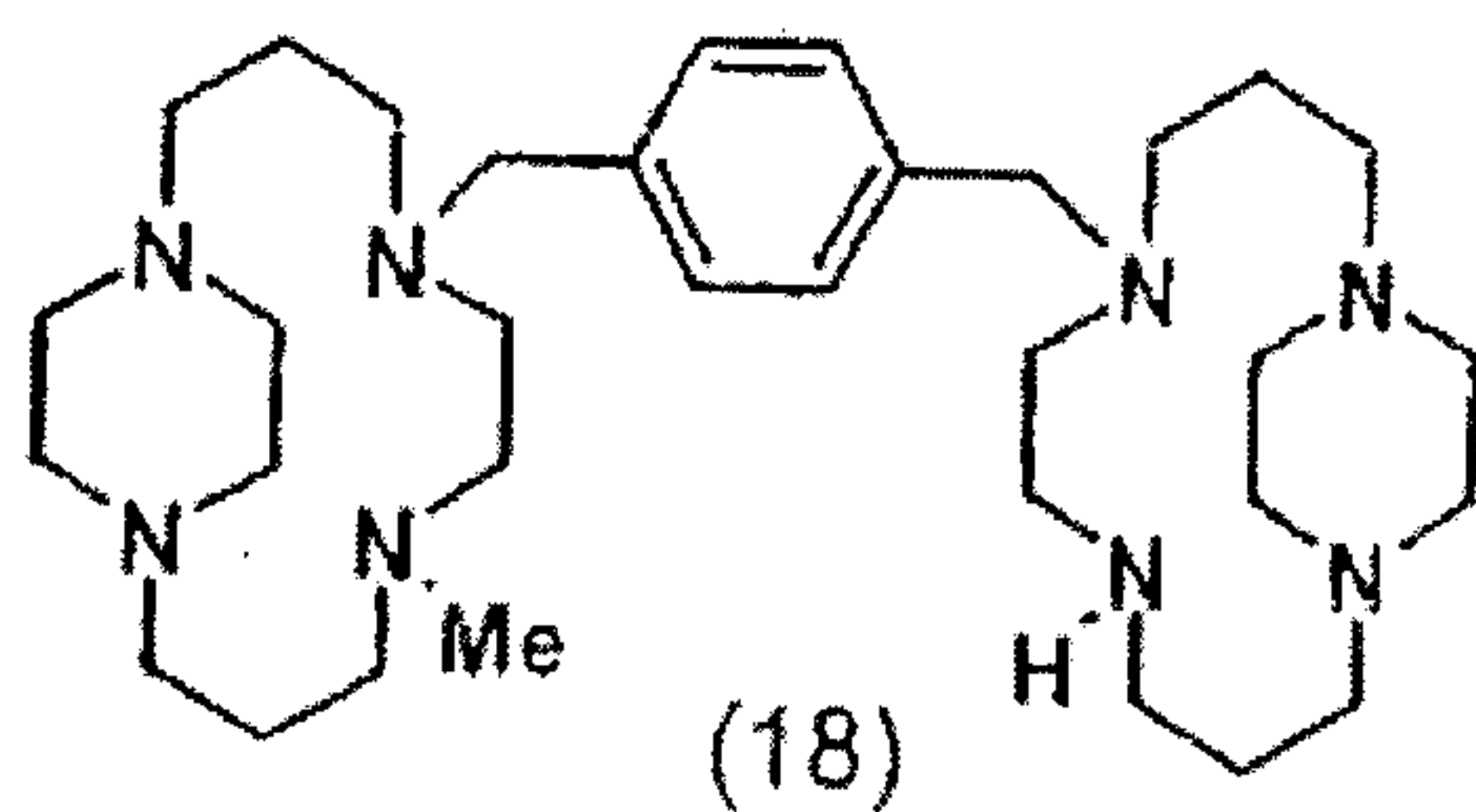
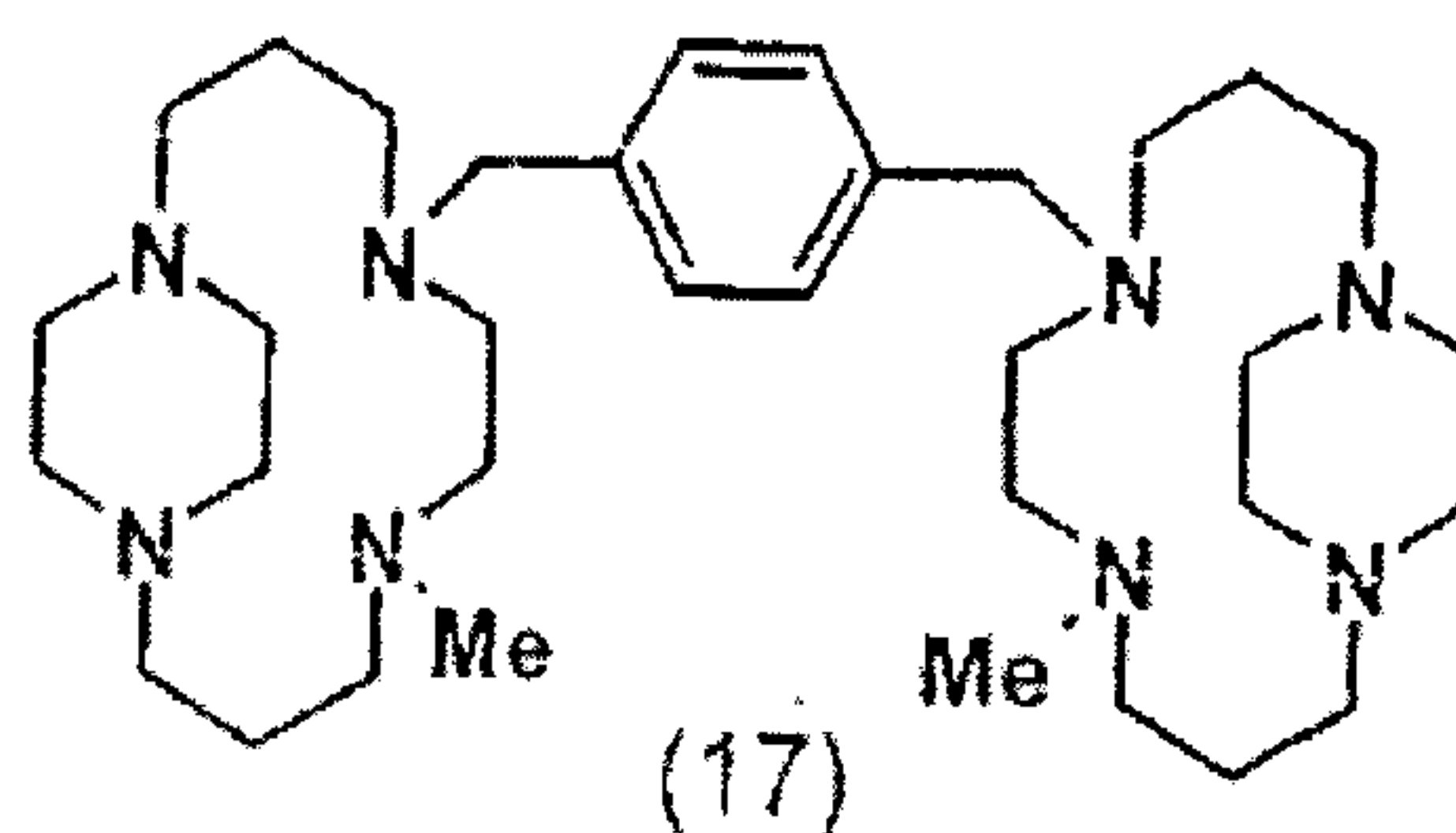
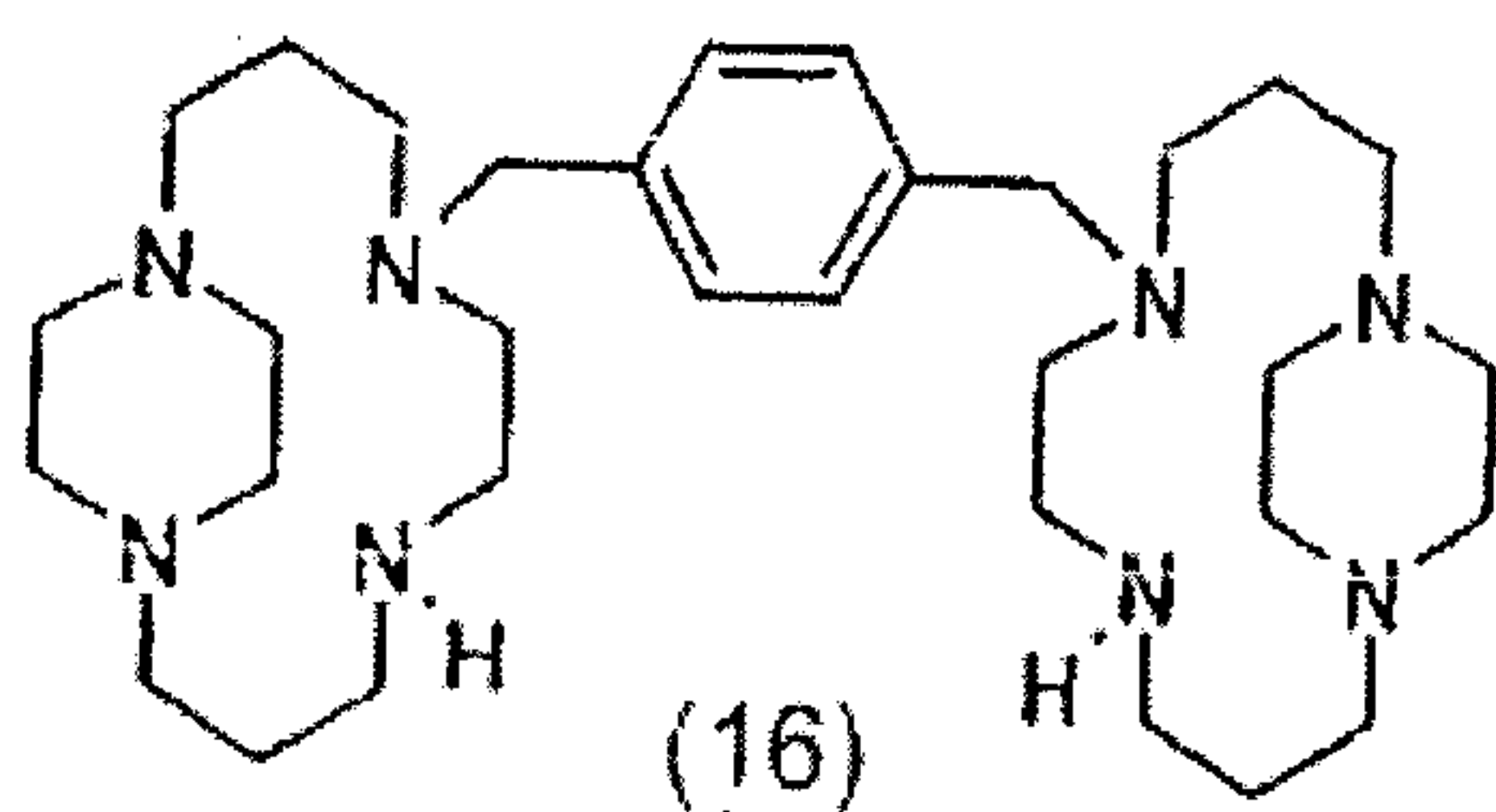
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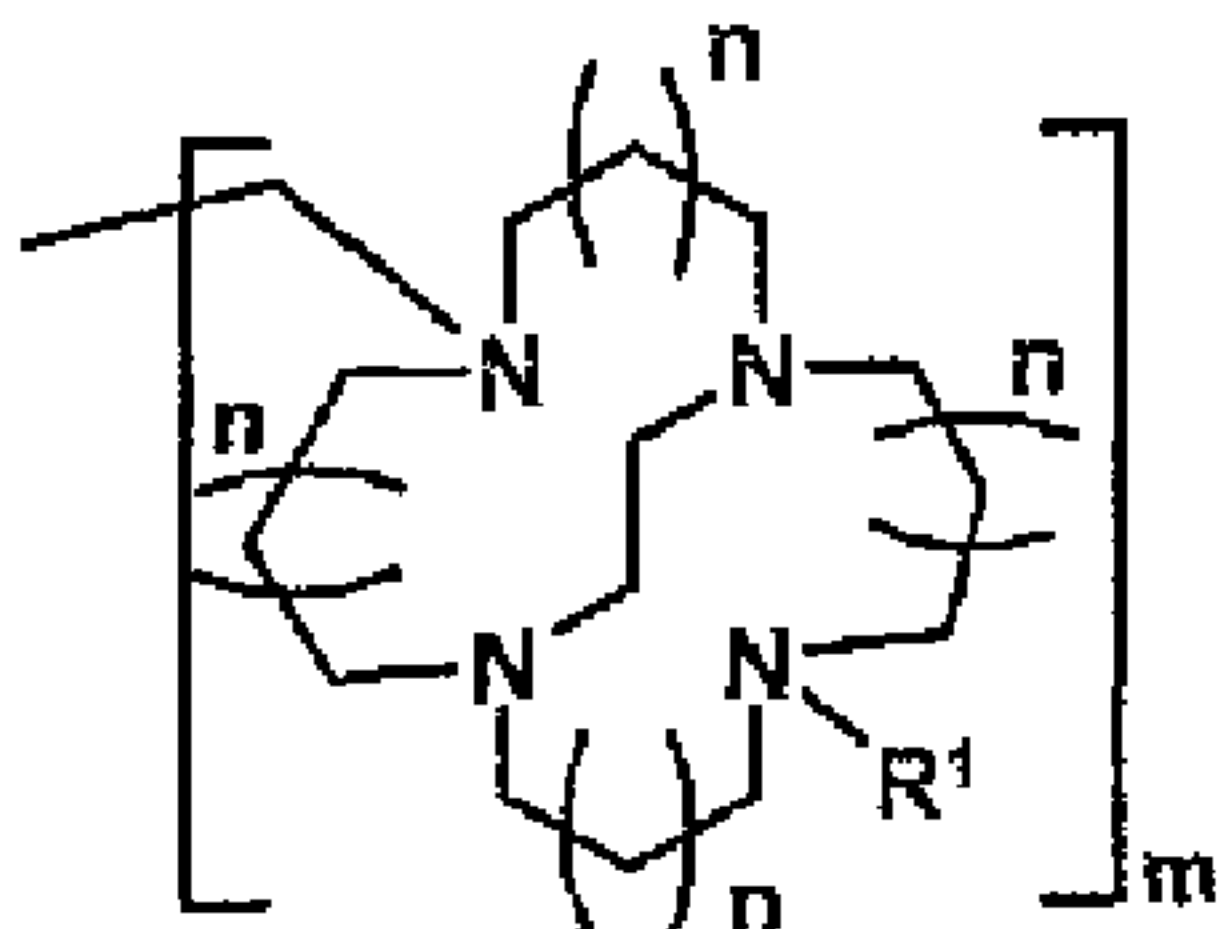
(15)

3. A compound as claimed in claim 1 or 2 wherein R is H or Me.
4. A compound as claimed in one of claims 1 to 3 wherein A_1 and A_2 are different.
5. A compound as claimed in any one of claims 1 to 4 complexed with a metal ion selected from transition metal elements and mixtures thereof.
6. A compound as claimed in any one of claims 1 to 5 labelled with a radioactive metal species or a fluorescent tag.

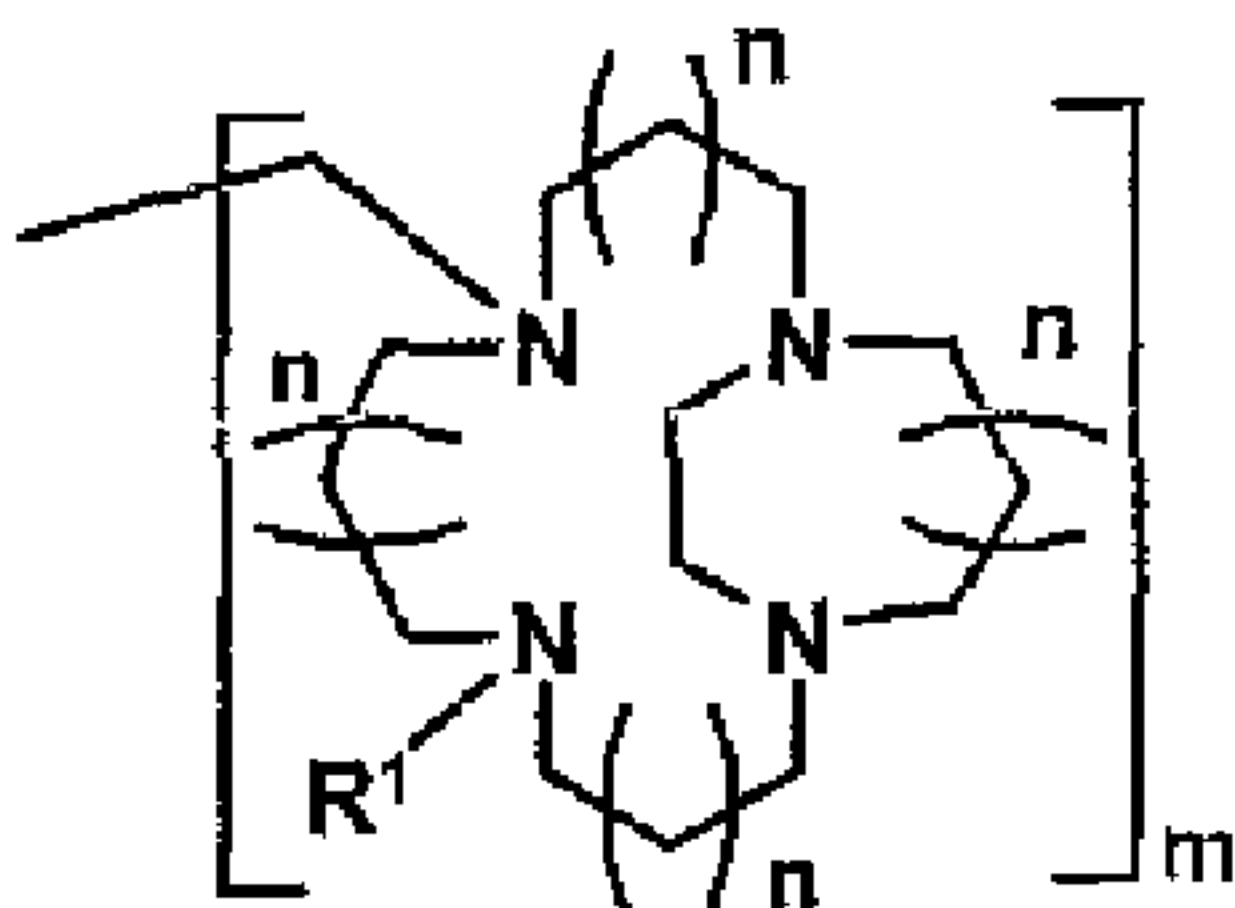
7. A compound as claimed in any one of claims 1 to 6 in which is inserted one or more radioisotopes.
8. A compound of formulae 16 to 23:



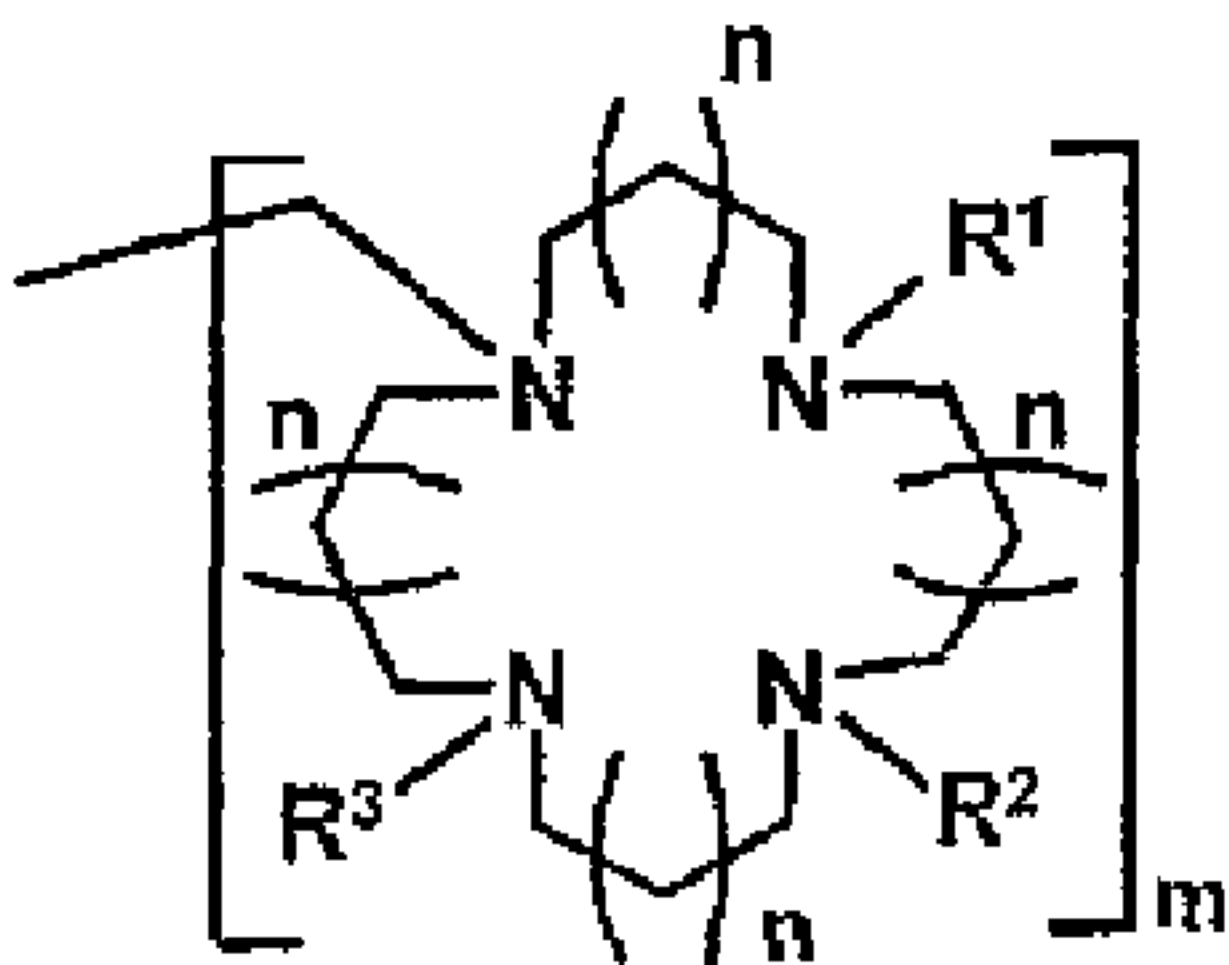
9. The use of a compound as claimed in any one of claims 1 to 8 as a CXCR4 co-receptor antagonist.
10. The use of a compound as claimed in any one of claims 1 to 8 for manufacture of an anti-viral dosage form.
11. The use of any compound as claimed in claim 10 for manufacture of a dosage form for treatment of HIV.
12. A pharmaceutical composition comprising a compound as claimed in any one of claims 1 to 8 and a pharmaceutically inert carrier.
13. Use of any of the compounds as claimed in any one of claims 1 to 8 for manufacture of a radiopharmaceutical for imaging chemokine receptors.
14. The use of any of the compounds as claimed in any one of claims 1 to 8 with a fluorescent tag for use in cellular or tissue imaging of the CXCR4 receptor.
15. The use of fluorescently labelled compounds claimed in drug screening assays for chemokine receptor binding.



(4)



(5)



(6)