CATALYSTS AND METHODS FOR POLYMER SYNTHESIS

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Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 46 days.

This patent is subject to a terminal disclaimer.

Appl. No.: 13/755,126

Filed: Jan. 31, 2013

Prior Publication Data


Related U.S. Application Data

Division of application No. 13/059,967, filed as application No. PCT/US2009/054773 on Aug. 24, 2009, now Pat. No. 8,633,123.

Provisional application No. 61/091,013, filed on Aug. 22, 2008, provisional application No. 61/096,313, filed on Sep. 11, 2008, provisional application No. 61/098,739, filed on Sep. 19, 2008.

Int. Cl.

B01J 31/00 (2006.01)
C07F 15/06 (2006.01)
C07F 37/18 (2006.01)
C07C 25/12 (2006.01)
C07C 48/704 (2006.01)
C07D 48/722 (2006.01)
C07F 9/535 (2006.01)
C07F 9/6561 (2006.01)
C08G 64/34 (2006.01)
C08F 4/80 (2006.01)
C07H 10/00 (2006.01)
B01J 37/00 (2006.01)
B01J 31/02 (2006.01)
B01J 31/22 (2006.01)

U.S. Cl.

CPC .......... C07F 15/065 (2013.01); B01J 31/182 (2013.01); C07C 25/124 (2013.01); C07C 279/12 (2013.01); C07D 213/53 (2013.01); C07D 48/704 (2013.01); C07D 48/722 (2013.01); C07F 9/535 (2013.01); C07F 9/6561 (2013.01); C08G 64/34 (2013.01); C08F 4/80 (2013.01); C07H 19/005 (2013.01);

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U.S. Appl. No. 13/755,112 claims *

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ABSTRACT

The present invention provides unimolecular metal complexes having increased activity in the copolymerization of carbon dioxide and epoxides. Also provided are methods of using such metal complexes in the synthesis of polymers. According to one aspect, the present invention provides metal complexes comprising an activating species with co-catalytic activity tethered to a multidentate ligand that is coordinated to the active metal center of the complex.

42 Claims, No Drawings
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CATALYSTS AND METHODS FOR POLYMER SYNTHESIS

PRIORITY CLAIM


BACKGROUND

Catalysts capable of effecting the copolymerization of epoxides and carbon dioxide to form aliphatic polycarbonates (APCs) have been known in the art since the 1960s. The early catalysts were based on heterogeneous zinc compounds and suffered from low reactivity, a lack of selectivity for polymer formation vs. cyclic carbonate formation, and a tendency to produce polycarbonates contaminated with ether linkages.

Improved catalysts based on transition metals have been discovered over the past decade or so. These newer catalysts have increased reactivity and improved selectivity. Nevertheless, even using highly active catalysts such as those disclosed in U.S. Pat. No. 7,304,172, the reaction times required to make high molecular weight polymer are typically quite long. In addition, the best-performing catalysts disclosed in the '172 patent require the addition of a separate co-catalyst to achieve optimum activity.

Attempts to address these shortcomings have been made. Catalysts described by Nozaki and co-workers (Angew. Chem. Int. Ed. 2006, 45, 7274-7277) tether an amine co-catalyst to a ligand of the catalyst. These next-generation catalytic systems suffer from lengthy and complicated syntheses and undesirable induction times prior to onset of polymerization. There remains a need for catalysts that have increased activity that will further reduce the polymerization time required to produce high molecular weight APCs.

SUMMARY

The present invention provides, among other things, unimolecular catalyst systems having activity in the copolymerization of carbon dioxide and epoxides and methods of using the same. In some embodiments, the present invention provides metal complexes having an activating species with co-catalytic activity tethered to a multidentate ligand that is coordinated to an active metal center of a metal complex.

In certain embodiments, the present invention provides unimolecular metal complexes and methods for using the same in the copolymerization of carbon dioxide and epoxides. In some embodiments, provided metal complexes have the structure:
or a combination of two or more of these,
wherein:
each occurrence of \( R^1 \) and \( R^2 \) is independently hydrogen or an optionally substituted radical selected from the group consisting of \( C_{1-20} \) aliphatic; \( C_{1-20} \) heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein \( R^1 \) and \( R^2 \) groups can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more additional heteroatoms;
each occurrence of \( R^3 \) is independently hydrogen or an optionally substituted radical selected from the group consisting of \( C_{1-20} \) aliphatic; \( C_{1-20} \) heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein an \( R^3 \) group can be taken with an \( R^1 \) or \( R^2 \) group to form one or more optionally substituted rings; and
each occurrence of \( R^4 \) is independently hydrogen, a hydroxyl protecting group, or an optionally substituted radical selected from the group consisting of \( C_{1-20} \) acyl; \( C_{1-20} \) aliphatic; \( C_{1-20} \) heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

In some embodiments, tethered activating functional groups \((Z)\) are cationic moieties. In certain embodiments, cationic moieties are selected from the group consisting of:
or a combination of two or more of these,

wherein:

each occurrence of R, R, and R is independently hydrogen or an optionally substituted radical selected from the group consisting of C_{1-20} aliphatic; C_{1-20} heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroary l ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 5- to 6-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heteroe cycl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein any two or more R, R, and R groups can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more additional heteroatoms;

R is hydrogen or —OR;

R is hydrogen, hydroxyl, or optionally substituted C_{1-20} aliphatic;

each occurrence of R and R is independently hydrogen or an optionally substituted radical selected from the group consisting of C_{1-20} aliphatic; C_{1-20} heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein R and R can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more heteroatoms, and an R or R group can be taken with an R or R group to form one or more optionally substituted rings;

each occurrence of R is independently hydrogen, a hydroxyl protecting group, or an optionally substituted radical selected from the group consisting of C_{1-20} acyl; C_{1-20} aliphatic; C_{1-20} heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or a combination of two or more of these,

each occurrence of R, R, and R is independently hydrogen or an optionally substituted radical selected from the group consisting of C_{1-20} aliphatic; C_{1-20} heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or a combination of two or more of these.
cyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein any two or more R², R⁷ and R⁸ groups can be taken together with intervening atoms to form one or more optionally substituted rings;
each occurrence of R¹₁ is independently selected from the group consisting of: halogen, —NO₂, —CN, —SR¹, —S(O)R², —S(O)₂R², —NR²(C(=O)R²), —OC(=O)R², —CO₂R², —NCO, —N₃, —OR², —OC(=O)NR², —N(R²)₂, —NR²C(O)R², —NR²C(O)OR² or an optionally substituted radical selected from the group consisting of C₁₋₅ aliphatic; C₁₋₅ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and a 1-5 heteroatom ring containing 0 to 4 heteroatoms.

X⁻ is any anion, and

Ring A is an optionally substituted, 5- to 10-membered heterocyclic group.

In some embodiments, an activating functional group (Z) is a phosphorus-containing functional group.

In certain embodiments, a phosphorus-containing functional group is chosen from the group consisting of: phosphines (—PR²); Phosphine oxides —P(O)R²; phosphinites P(OR²)₂; phosphinites P(OR²)²; phosphates OP(OR²)₂; phosphates PO(OR²)₂; phosphonites OP(OR²)₂; phosphonites PO(OR²)₂; phosphonium salts ([—PR²]⁺); where the phosphorus-containing functional group may be linked to a metal complex through any available position (e.g., direct linkage via the phosphorus atom, or in some cases via an oxygen atom).

In certain embodiments, a phosphorus-containing functional group is chosen from the group consisting of:

- \((X)_{b} - ([R²(=O)R³]_a)_{b}Q^{n}\), wherein:
  - X is —O—, —N—, or —NR²—,
  - \(b\) is 1 or 0,
  - each of R², R⁷ and R⁸ is independently present or absent and, if present, are independently selected from the group consisting of optionally substituted C₁₋₅ aliphatic, optionally substituted C₆₋₁₄ aryl, optionally substituted 3- to 14-membered heterocyclic, optionally substituted 5- to 14-membered heteroaryl, halogen, —OR², —OR³, —NR², and N(R²)₂ where R² is hydrogen, or an optionally substituted C₁₋₅ aliphatic, optionally substituted 6- to
14-membered aryl, optionally substituted 3- to 14-membered heterocyclic, or optionally substituted 5- to 14-membered heterocyclic, Q is any anion, and n is an integer between 1 and 4.

In some embodiments, the present disclosure encompasses methods for the copolymerization of epoxides and carbon dioxide, such methods comprising contacting one or more epoxides with a catalyst described above in the presence of carbon dioxide.

In some embodiments, the present disclosure encompasses methods for the formation of cyclic carbonates from epoxides and carbon dioxide, such methods comprising contacting one or more epoxides with a catalyst described above in the presence of carbon dioxide.

In some embodiments, the present disclosure encompasses methods for the formation of polymers, such methods comprising contacting one or more epoxides with a catalyst described above.

DEFINITIONS

Definitions of specific functional groups and chemical terms are described in more detail below. For purposes of this invention, the chemical elements are identified in accordance with the Periodic Table of the Elements, CAS version, Handbook of Chemistry and Physics, 75th Ed., inside cover, and specific functional groups are generally defined as described therein. Additionally, general principles of organic chemistry, as well as specific functional moieties and reactivity, are described in Organic Chemistry, Thomas Sorrell, University Science Books, Sausalito, 1999; Smith and March’s Advanced Organic Chemistry, 5th Edition, John Wiley & Sons, Inc., New York, 2001; LaRoc, Comprehensive Organic Transformations, VCH Publishers, Inc., New York, 1989; Curranthers, Some Modern Methods of Organic Synthesis, 3rd Edition, Cambridge University Press, Cambridge, 1987; the entire contents of each of which are incorporated herein by reference.

Certain compounds of the present invention can comprise one or more asymmetric centers, and thus can exist in various stereoisomeric forms, e.g., enantiomers and/or diastereomers. Thus, inventive compounds and compositions thereof may be in the form of an individual enantiomer, diastereomer or geometric isomer, or may be in the form of a mixture of stereoisomers. In certain embodiments, the compounds of the invention are enantiopure compounds. In certain embodiments, mixtures of enantiomers or diastereomers are provided.

Furthermore, certain compounds, as described herein may have one or more double bonds that can exist as either a Z or E isomer, unless otherwise indicated. The invention additionally encompasses the compounds as individual isomers substantially free of other isomers and alternatively, as mixtures of various isomers, e.g., racemic mixtures of enantiomers. In addition to the above-mentioned compounds per se, this invention also encompasses compositions comprising one or more compounds.

As used herein, the term “isomers” includes any and all geometric isomers and stereoisomers. For example, “isomers” include cis- and trans-isomers, E- and Z-isomers, R- and S-enantiomers, diastereomers, (o)-isomers, (t)-isomers, racemic mixtures thereof, and other mixtures thereof, as falling within the scope of the invention. For instance, a compound may, in some embodiments, be provided substantially free of one or more corresponding stereoisomers, and may also be referred to as “stereocchemically enriched.”

Where a particular enantiomer is preferred, it may, in some embodiments be provided substantially free of the opposite enantiomer, and may also be referred to as “optically enriched.” “Optically enriched,” as used herein, means that the compound is made up of a significantly greater proportion of one enantiomer. In certain embodiments the compound is made up of at least about 90% by weight of an enantiomer. In some embodiments the compound is made up of at least about 95%, 97%, 98%, 99%, 99.5%, 99.7%, 99.8%, or 99.9% by weight of an enantiomer. In some embodiments the enantiomeric excess of provided compounds is at least about 90%, 95%, 97%, 98%, 99%, 99.5%, 99.7%, 99.8%, or 99.9%. In some embodiments, enantiomers may be isolated from racemic mixtures by any method known to those skilled in the art, including chiral high pressure liquid chromatography (HPLC) and the formation and crystallization of chiral salts or prepared by asymmetric syntheses. See, for example, Jacques, et al., Enantiomers, Racemates and Resolutions (Wiley Interscience, New York, 1981); Wilen, S. H., et al., Tetrahedron 33:2725 (1977); Eliel, E. L. Stereochemistry of Carbon Compounds (McGraw-Hill, NY, 1962); Wilen, S. H., Tables of Resolving Agents and Optical Resolutions p. 268 (E. L. Eliel, Ed., Univ. of Notre Dame Press, Notre Dame, Ind. 1972).

The terms “halo” and “halogen” as used herein refer to an atom selected from fluorine (fluoro, —F), chlorine (chloro, —Cl), bromine (bromo, —Br), and iodine (iodo, —I).

The term “aliphatic” or “aliphatic group”, as used herein, denotes a hydrocarbon moiety that may be straight-chain (i.e., unbranched), branched, or cyclic (including fused, bridging, and spiro-fused polycyclic) and may be completely saturated or may contain one or more units of unsaturation, but which is not aromatic. Unless otherwise specified, aliphatic groups contain 1-30 carbon atoms. In certain embodiments, aliphatic groups contain 1-12 carbon atoms. In certain embodiments, aliphatic groups contain 1-8 carbon atoms. In certain embodiments, aliphatic groups contain 1-6 carbon atoms. In some embodiments, aliphatic groups contain 1-5 carbon atoms, in some embodiments, aliphatic groups contain 1-4 carbon atoms, in yet other embodiments aliphatic groups contain 1-3 carbon atoms, and in yet other embodiments aliphatic groups contain 1-2 carbon atoms. Suitable aliphatic groups include, but are not limited to, linear or branched, alkyl, alkenyl, and alkynyl groups, and hybrids thereof such as (cyloalkyl)alkyl, (cyloalkenyl)alkyl or (cyloalkynyl)alkenyl. In certain embodiments, the term aliphatic group encompasses aliphatic groups wherein one or more hydrogen atoms are replaced with a halogen atom. In certain embodiments, the term aliphatic group encompasses chlorinated or fluorinated aliphatic groups including perborinated compounds.

The term “epoxide”, as used herein, refers to a substituted or unsubstituted oxirane. Such substituted oxiranes include monosubstituted oxiranes, disubstituted oxiranes, trisubstituted oxiranes, and tetrasubstituted oxiranes. Such epoxides may be further optionally substituted as defined herein. In certain embodiments, epoxides comprise a single oxirane moiety. In certain embodiments, epoxides comprise two or more oxirane moieties.

The term “polymer”, as used herein, refers to a molecule of high relative molecular mass, the structure of which comprises the multiple repetition of units derived, actually or conceptually, from molecules of low relative molecular mass. In certain embodiments, a polymer is comprised of only one monomer species (e.g., polyethylene oxide). In certain embodiments, a polymer of the present invention is a copolymer, terpolymer, heteropolymer, block copolymer, or tapered heteropolymer of one or more epoxides.
The term "unsaturated", as used herein, means that a moiety has one or more double or triple bonds.

The terms "cyclolipaphatic", "carbocycle", or "carbocyclic", used alone or as part of a larger moiety, refer to a saturated or partially unsaturated cyclic aliphatic monomeric, bicyclic, or polycyclic ring systems, as described herein, having from 3 to 12 members, wherein the aliphatic ring system is optionally substituted as defined above and described herein. Cyclolipaphatic groups include, without limitation, cyclopropyl, cyclobutyl, cyclopentyl, cyclopentenyl, cyclohexyl, cyclohexenyl, cycloheptyl, cycloheptenyl, cyclooctyl, cyclooctenyl, and cyclooctatetraenyl. In some embodiments, the cycloalkyl has 3-6 carbons. The terms "cyclolipaphatic", "carbocycle" or "carbocyclic" also include aliphatic groups that are fused to one or more aromatic or nonaromatic rings, such as decalynylphiphyl or tetralinylphiphyl, where the radical or point of attachment is on the aliphatic ring. In some embodiments, a carbocyclic groups is bicyclic. In some embodiments, a carbocyclic group is tricyclic. In some embodiments, a carbocyclic group is polycyclic.

The term "alkyl", as used herein, refers to saturated, straight- or branched-chain hydrocarbon radicals derived by removal of a single hydrogen atom from an aliphatic moiety. Unless otherwise specified, alkyl groups contain 1-12 carbon atoms. In certain embodiments, alkyl groups contain 1-8 carbon atoms. In certain embodiments, alkyl groups contain 1-6 carbon atoms. In some embodiments, alkyl groups contain 1-5 carbon atoms, in some embodiments, alkyl groups contain 1-4 carbon atoms, in yet other embodiments alkyl groups contain 1-3 carbon atoms, and in yet other embodiments alkyl groups contain 1-2 carbon atoms. Examples of alkyl radicals include, but are not limited to, methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, sec-butyl, sec-pentyl, iso-pentyl, tert-butyl, n-pentyl, neopentyl, n-hexyl, sec-hexyl, n-heptyl, n-octyl, n-decyl, n-undecyl, dodecyl, and the like.

The term "alkenyl" as used herein, denotes a monovalent group derived by the removal of a single hydrogen atom from a straight- or branched-chain aliphatic moiety having at least one carbon-carbon double bond. Unless otherwise specified, alkenyl groups contain 2-12 carbon atoms. In certain embodiments, alkenyl groups contain 2-8 carbon atoms. In certain embodiments, alkenyl groups contain 2-6 carbon atoms. In some embodiments, alkenyl groups contain 2-5 carbon atoms, in some embodiments, alkenyl groups contain 2-4 carbon atoms, in yet other embodiments alkenyl groups contain 2-3 carbon atoms, and in yet other embodiments alkenyl groups contain 2 carbon atoms. Alkenyl groups include, for example, ethenyl, propenyl, allyl, 1,3-butadienyl, butenyl, 1-methyl-2-buten-1-yl, allyl, 1,3-butadienyl, alleny, and the like.

The term "alkynyl", as used herein, refers to a monovalent group derived by the removal of a single hydrogen atom from a straight- or branched-chain aliphatic moiety having at least one carbon-carbon triple bond. Unless otherwise specified, alkynyl groups contain 2-12 carbon atoms. In certain embodiments, alkynyl groups contain 2-8 carbon atoms. In certain embodiments, alkynyl groups contain 2-6 carbon atoms. In some embodiments, alkynyl groups contain 2-5 carbon atoms, in some embodiments, alkynyl groups contain 2-4 carbon atoms, in yet other embodiments alkynyl groups contain 2-3 carbon atoms, and in yet other embodiments alkynyl groups contain 2 carbon atoms. Representative alkynyl groups include, but are not limited to, ethynyl, 2-propynyl (propargyl), 1-propynyl, and the like.

The term "carbocycle" and "carbocyclic ring" as used herein, refer to monocyclic and polycyclic moieties wherein the rings contain only carbon atoms. Unless otherwise specified, carbocycles may be saturated, partially unsaturated or aromatic, and contain 3 to 20 carbon atoms. Representative carbocycles include cyclopropane, cyclobutane, cyclopentane, cyclohexane, bicyclo[2.2.2]octane, norbornane, phenyl, cyclohexene, naphthalene, and spiro[4.5]decanne, to name but a few.

The term "aryl" used alone or as part of a larger moiety as in "aryalkyl", "arylalkoxy", or "aryloxyalkyl", refers to monocyclic and polycyclic ring systems having a total of six to 20 ring members, wherein at least one ring in the system is aromatic and wherein each ring in the system contains three to twelve ring members. The term "aryl" may be used interchangeably with the term "aryl ring". In certain embodiments of the present invention, "aryl" refers to an aromatic ring system which includes, but is not limited to, phenyl, biphenyl, naphthyl, anthracenyl and the like, which may bear one or more substituents. Also included within the scope of the term "aryl", as it is used herein, is a group in which an aromatic ring is fused to one or more additional rings, such as benzofernalyl, indanlyl, phthalimidyl, naphthimidyl, phenantrimidyl, or tetrahydrodiphosphyl, and the like.

The term "hetaryl", as used herein, refers to aliphatic groups wherein one or more carbon atoms are independently replaced by one or more atoms selected from the group consisting of oxygen, sulfur, nitrogen, phosphorus, or boron. In certain embodiments, one or six carbon atoms are independently replaced by one or more of oxygen, sulfur, nitrogen, or phosphorus. Heteroaliphatic groups may be substituted or unsubstituted, branched or unbranched, cyclic or acyclic, and include saturated, unsaturated or partially unsaturated groups.

The terms "heteroaryl" and "heteroar", used alone or as part of a larger moiety, e.g., "hetoraryalkyl", or "heteroarylalkoxy", refer to groups having 5 to 14 ring atoms, preferably 5, 6, or 9 ring atoms; having 6, 10, or 14 or more electrons shared in a cyclic array; and having, in addition to carbon atoms, from one to five heteroatoms. The term "heteroaromat" refers to nitrogen, oxygen, or sulfur, and includes any oxidized form of nitrogen or sulfur, or any quaternized form of a basic nitrogen. Heteroaliphatic groups include, without limitation, thienyl, furanyl, pyrrolyl, imidazolyl, pyrazolyl, triazolyl, tetrazolyl, oxazolyl, isoxazolyl, oxadiazolyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyridazinyl, pyrimidinyl, pyrazinyl, indolizinyl, purinyl, naphthyridinyl, benzofuranyl and pteridinyl. The terms "heteroary" and "heteroar"-, as used herein, also include groups in which a heteroaromatic ring is fused to one or more ary, cyclolipaphatic, or heterocyclic rings, where the radical or point of attachment is on the heteroaromatic ring. Nonlimiting examples include indolyl, isoindolyl, benzothienyl, benzofuranyl, dibenzofuranyl, indazolyl, benzimidazolyl, benzthiazolyl, quinolyl, isoquinolyl, cinnolinyl, phthalazinyl, quinoxalinyl, 4H-quinolinyl, carbazolyl, acridinyl, phenazinyl, phenothiazinyl, phenoazinyl, tetrahydroisoquinolinyl, and pyridino[2,3-b]1,4-oxazin-3 (4H)-one. A heteroaryl group may be mono- or polycyclic. The term "heteroary" may be used interchangeably with the terms "heteroaryl ring", "heteroaryl group", or "heteroaromatic", any of which terms include rings that are optionally substituted. The term "hetoraryalkyl" refers to an alkyl group substituted by a heteroaryl, wherein the alkyl and heteroaryl portions independently are optionally substituted.

As used herein, the terms "heterocycle", "heterocycl", "heterocyclic radical", and "heterocyclic ring" are used interchangeably and refer to a stable 5- to 7-membered monocyclic or 7-14-membered bicyclic heterocyclic moiety that is saturated, partially unsaturated, or aromatic and having, in
addition to carbon atoms, one or more, preferably one to four, heteroatoms, as defined above. When used in reference to a ring atom of a heterocycle, the term “nitrogen” includes a substituted nitrogen. As an example, in a saturated or partially unsaturated ring having 0-3 heteroatoms selected from oxygen, sulfur or nitrogen, the nitrogen may be N (as in 3,4-dihydro-2H-pyrrolyl), NH (as in pyrrolidinyl), or NR (as in N-substituted pyrrolidinyl).

A heterocyclic ring can be attached to its pendant group at any heteroatom or carbon atom that results in a stable structure and any of the ring atoms can be optionally substituted. Examples of such saturated or partially unsaturated heterocyclic radicals include, without limitation, tetrahydrofuranyl, tetrahydrothiophenyl, pyrrolidinyl, piperidinyl, pyrrolinyl, tetrahydroquinolinyl, tetrahydroisoquinolinyl, decahydroquinolinyl, oxazolidinyl, pipеразинyl, dioxanyl, dioxolany, diazepinyl, oxazepinyl, diazepinyl, morpholinyl, and quinuclidinyl. The terms “heterocyclic,” “heterocyclyl,” “heterocyclyl ring,” “heterocyclic group,” “heterocyclic moiety,” and “heterocyclic radical,” are used interchangeably herein, and also include groups in which a heterocyclyl ring is fused to one or more aryl, heteroaryl, or cycloalkyl radicals, such as indolyl, 3H-indolyl, chromanyl, phenantridinyl, or tetrahydroquinolinyl, where the radical or point of attachment is on the heterocyclyl ring. A heterocyclyl group may be mono- or bicyclic. The term “heterocyclylalkyl” refers to an alkyl group substituted by a heterocyclyl, wherein the alkyl and heterocyclyl portions independently are optionally substituted.

The term “acyl” as used herein refers to a group having a formula —C(=O)R where R is hydrogen or an optionally substituted aliphatic, aryl, or heterocyclyl group.

As used herein, the term “partially unsaturated” refers to a ring moiety that includes at least one double or triple bond. The term “partially unsaturated” is intended to encompass rings having multiple sites of unsaturation, but not intended to include aryl or heteroaryl moieties, as herein defined. One of ordinary skill in the art will appreciate that the synthetic methods, as described herein, utilize a variety of protecting groups. By the term “protecting group,” as used herein, it is meant that a particular functional moiety, e.g., O, S, or N, is masked or blocked, permitting, if desired, a reaction to be carried out selectively at another reactive site in a multifunctional compound. In preferred embodiments, a protecting group reacts selectively in good yield to give a protected substrate that is stable to the projected reactions; the protecting group is preferably selectively removable by readily available, preferably non-toxic reagents that do not attack the other functional groups; the protecting group forms a separable derivative (more preferably without the generation of new sterogenic centers); and the protecting group will preferably have a minimum of additional functionality to avoid further sites of reaction. As detailed herein, oxygen, sulfur, nitrogen, and carbon protecting groups may be utilized. By way of non-limiting example, hydroxyl protecting groups include methyl, methoxymethyl (MOM), MOMethylthiomethyl (MTM), t-butylthiomethyl, (phenyldimethylsilyl) methoxymethyl (SMOM), benzoxymethyl (BOM), p-methoxybenzoxymethyl (PMBM), (4-methoxyphenoxymethyl (p-AOM), guaiacolmethyl (GUM), t-butoxymethyl, 4-pentenylmethyl (POM), siloxymethyl, 2-methoxyethoxymethyl (MEM), 2,2,2-trichloroethoxymethyl, bis(2-chloroethoxymethyl, 2-(trimethyloxysilyl)ethoxymethyl (SEOM), tetrahydropropyl (TTH), 3-bromotetrahydropropyranyl, tetrahydropropyl (TTH), 1-methoxycyclohexyl, 4-methoxytetrahydropropyranyl (MTHP), 4-methoxytetrahydropropyranyl, 4-methoxytetrahydropropyranyl S,S-dioxide, 1-(4-chloro-4-methylphenyl)-1-methoxypiperidin-4-yl (CTMP), 1,4-dioxan-2-yl, tetrahydrofuranyl, tetrahydrofuranoyl, 2,3,3a,4,5,6,7a-octahydro-7,8,8-trimethyl-4,7-methanothieno[2,3-b]furanyl, 1-ethoxyethyl, 1-(2-chlorothioethyl, 1-methyl-1-ethoxymethyl, 1-methyl-1-benzoxylxy-2-thioxoethyl, 2,2-trichloromethyl, 2-trimethylisobutyryl, 2-(phenylisocyanato)ethyl, tert-butyl, allyl, p-chlorophenyl, p-methoxyphenyl, 2,4-dinitrophenyl, benzyl, p-methoxybenzyl, 3,4-dimethoxybenzyl, o-nitrobenzyl, p-nitrobenzyl, p-halobenzyl, 2,6-dichlorobenzyl, p-cyanobenzyl, p-phenylbenzyl, 2-picoly, 4-picolly, 3-methyl-2-picyl N-oxide, diphenylmethyl, p,p'-dinitrobenzyl, 5-dibenzosuberyl, triphenylmethy1, α-naphthylpheny1methyl, p-methoxyphenylphenylmethyl, diphenylethylmethyl, 4,4,4'-tris(4,4-dichlorophenacyl)phenylmethyl, 4,4,4'-tris(benzoyloxyphenyl)alkyl, 3-(imidazol-1-yl)bis(4,4-dimethylphenyl)methyl, 1,1-bis(4-methylphenyl)-1-pyrylenylmethyl, 9-antryl, 9-(phenylxanthenyl, 9-phenyl-10-oxo)anthryl, 1,3-benzodiazenyl, benzosathiokyl S,S-dioxide, trimethylsilyl (TMS), triethylsilyl (TES), trimethylsilyl (TIPS), dimethylsilylpropyl (DPM), diethylisopropylsilyl (DEIPS), dimethyldimethylsilyl, t-butyl(dimethylsilyl)TBDS, t-butylphenylsilyl (TBP), tribenzylsilyl, tri-p-xylisilyl, triphenylsilyl, diphenylmethyl (DPM), t-butylmethoxyphenylsilyl (TBMP), formate, benzoylformate, acetate, chloroacetate, dichloroacetate, trichloroacetate, trifluoroacetate, methoxycetate, triphenyloxycetate, phenoxycetate, p-chlorophenoxyacetate, 3-phenylpropionate, 4-oxopentanoate (levulinate), 4,4-(ethylidenedithio)pentanoate (levulinoylthiobaccatol), pivaloate, adamanatoate, eronate, 4-methycrotonate, benzate, p-phenylbenzoate, 2,4,6-trimethylbenzoate (mesitato), alkyl methyl carbonate, 9-fluorenylmethyl carbonate (Fmoc), alkyl ethyl carbonate, alkyl 2,2,2-trichloroethyl carbonate (Troc), 2-(trimethylsilyl)ethy1 carbonate (TMSEC), 2-(phenylsulfonyl)ethyl carbonate (Psec), 2-(triphenylphosphinothio)ethyl carbonate (Psec), alkyl isobutyrate, alkyl vinyl carbonate, alkyl alkyl carbonate, alkyl p-nitrophenyl carbonate, alkyl benzyl carbonate, alkyl p-methoxybenzyl carbonate, alkyl 3,4-dimethoxybenzyl carbonate, alkyl o-nitrobenzyl carbonate, alkyl p-nitrobenzyl carbonate, alkyl S-benzyl thiocarbonate, 4-ethoxy-1-naphthyl carbonate, methyl thioisocyanate, 2,4-dichlorobenzene, 4-azidoxybutyrace, 4-nitro-4-methylpentanoate, o-(dibromomethyl) benzoxate, 2-formylbenzenesulfonate, 2-(methylthio)methoxymethyl, 4-(methylthiomethoxy)butyrate, 2-(methylthiophenoxymethyl)benzoate, 2,2,2-dichloro-4-methylphenoxacetate, 2,6-dichloro-4-(1,1,3,3-tetramethylbutyl)phenoxacetate, 2,4-bis(1,1-dimethylpropyl)phenoxacetate, chloridiphenylacetate, isobutyrate, monosuccinate, (E)-2-methyl-2-butoenate, o-((methoxycarbonyl)benzoxate, α-naphthoate, nitrate, alkyl N,N,N,N'-tetramethylephosphorodiamidate, alkyl N-phenylcarbamate, borate, dimethylphosphonitriyl, alkyl 2,4-dinitrophenylsulfonate, sulfate, methanesulfonate (mesylate), benzyloximate, and tosylate (Ts). For protecting 1,2- or 1,3-diols, the protecting groups include methylene acetal, ethylenediic acetal, 1,1-butyldiethylidene ketal, 1-phenylthiolidene ketal, (4-methoxyphenyl)ethylenediic acetal, 2,2,2-trichlorothylidene acetal, acetonide, cyclopentaenedi ketal, cyclohexyldiene ketal, cycloheptydene ketal, benzylidene acetal, p-methoxybenzylidene acetal, 2,4-dimethoxybenzylidene ketal, 3,4-dimethoxybenzylidene acetal, 2-nitrobenzylidene acetal, methoxymethylene acetal, ethoxymethylene acetal,
dimethoxymethylene ortho ester, 1-methoxymethylene ortho ester, 1-ethoxymethylene ortho ester, 1,2-dimethoxymethylene ortho ester, α-methoxybenzylidene ortho ester, 1-(N-N-dimethylamino)methylene derivative, 1,3-(1,3,3,3-tetraoxopropylsiloxy)benzene derivative (TIPS), tetra-t-butoxydisiloxane-1,3-diyldiene derivative (TBDS), cyclic carbonates, cyclic boronates, ethyl boronate, and phenyl boronate. Amino-protecting groups include methyl carbamate, ethyl carbamate, 9-fluorenylmethyl carbamate (Fmoc), 9-(2-sulfo)fluorenylmethyl carbamate, 9-(2,7-dibromo)fluorene-
methyl carbamate, 2,7-di-4-butyryl[9-(10,10-dioxo-10, 10,10-tetrahydrothioxan oxyl)]methyl carbamate (BDD-Tmoc), 4-methoxyphenacyl carbamate (Phenoc), 2,2,2-trichloroethyl carbamate (Trcoc), 2-trimethylsilyl ethyl carbamate (Tcoc), 2-phénynyl ethyl carbamate (Hzc), 1-(1-adamantyl)-1-methyl ethyl carbamate (Adoc), 1,1-dimethyl-2-haloethyl carbamate, 1,1-dimethyl-2,2-dibromoethyl carbamate (DHBoc), 1,1-dimethyl-2,2-trichloroethyl carbamate (TCOC), 1,1-dimethyl(4-biphenylyl)ethyl carbamate (Bpoc), 1-(3,5-di-4-butyryl)-1-methyl ethyl carbamate (T-BnBoc), 2-[(4-pyridyl)ethoxy]methyl carbamate (Pyoc), 2-(N,N-dicyclohexylcarbamoyl)ethyl carbamate, t-butyl carbamate (Boc), 1-adamantyl carbamate (Adoc), vinyl carbamate (Voc), allyl carbamate (Alloc), iso-propenyl carbamate (Ipcoc), cinnamyl carbamate (Coc), 4-nitrocin narnyl carbamate (Nom), 2,8-quinolyl carbamate, N-hydroxysyipiperidinyl carbamate, allyldithio carbamate, benzyl carbamate (Cbz), p-methoxybenzyl carbamate (Moz), p-nitrobenzyl carbamate, p-bromobenzyl carbamate, p-chlorobenzyl carbamate, 2,4-dichlorobenzyl carbamate, p-methylsulfinyl benzyl carbamate (Msz), 9-anthrylmethyl carbamate, diphenylmethylen carbamate, 2-methylthioethyl carbamate, 2-methylsulfonyl ethyl carbamate, 2-(p-toluene sulfonyl)ethyl carbamate, 2-(1,3-dithianyl)methyl carbamate (Dmoc), 4-methylthiophenyl carbamate (Mtpc), 2,4-dimethylthiophenyl carbamate (Bmmpc), 2-phosphonomethyl carbamate (Ppeoc), 2-triphenylphosphonioisopropyl carbam ate (Ppoeoc), 1,1-dimethyl-2-cyanoethyl carbamate, m-chlorop-acycloxybenzyl carbamate, p-(di-hydroxobenzyl)benzyl carbamate, 5-benzoxazolylmethyl carbamate, 2-(trifluoromethyl)-6-chromon methyl carbamate (Tcrco), m-nitrophenyl carbamate, 3,5-dimethoxycarbonyl benzenediazonium tetrafluoroborate, 3,4-dimethoxy-6-nitrobenzyl carbamate, phenoxy[1-nitrophenyl]methyl carbamate, phenothiazinyl(+1)-carbonyl derivative, N'-p-toluenesulfonylamin ocarbonyl derivative, N'-phenyilmethiothiocarbonyl derivative, 1-methyl carbamate, 3-hydroxy diazocarbonyl carbamate, cyclobutyl carbamate, cyclohexyl carb amate, cyclopentyl carbamate, cyclopropylmethyl carbamate, 2,6-dimethoxy carbonylvinyl carbamate, o-(N,N-dimethylcarbamoid benzyl carbamate, 1,1-dimethyl-3 (N,N-dimethylcarbamido)propyl carbamate, 1,1-dimethyl-3-dimethyl propynyl carbamate, di(2-pyridyl) methyl carbamate, 2-fluoroethyl carbamate, 2-iodoethyl carbamate, isobornyl carbamate, isobutyl carbamate, isocionetyl carbamate, (p-(methoxyphenylazo)benzyl carbamate, 1-methylcyclobutyl carbamate, 1-methylcyclohexyl carbamate, 1-methyl-1-cyclopentylmethyl carbamate, 1-methyl-1-(3,5-dimethoxy)ethyl carbamate, 1-methyl-1-(p-phenylazophenyl)ethyl carbamate, 1-(methyl-1-phenylethyl) carbamate, 2,4,6-tri-4- butylphenyl carbamate, 4-(trimethylammonium) benzyl carbamate, 2,4,6-trimethylbenzyl carbamate, formamide, acetam ides, chloracetamide, trichloroacetamide, trifluoroacetamide, phenylacetamide, 3-phenylpropanamide, picolinamide, 3-pyridylcarboxamide, N-benzoylphenylalanine derivative, benzamide, p-phenylbenzamide, o-nitrophen yacetamide, o-nitrophenoxacyacetamide, acetacetamide, (N,N-di-thiobenzoylcarbimino)acetamide, (N-p-hydroxyphenyl)propanamide, (3-nitrophenyl)propanamide, 2-methyl-2-(o-nitrophenyl)propanamide, 2-methyl-2-(o-phenylazo)propanamide, 4-chlorobutanamide, 3-methyl-3-nitrobutanamide, o-nitrocinnamidine, N-acetylthionine derivative, o-nitrobenzamide, o-(benzoylom ethyl)benzamide, 4,5-diphenyl-3-oxazolin-2-one, N-phthal imide, N-dithiasuccinimide (Dts), N-2,3-diphenylmaleimide, N-2,5-dimethylpyrrole, N-1,1,4-tetramethyldisilazancelyclopentane adduct (STABase), 5-substituted 1,3-dimethyl-1,3,5-triazacyclohexan-2-one, 5-substituted 1,3-dimethyl-1,3,5-triazacyclohexan-2-one, 5-substituted 3,5-dinitro-4-pyridone, N-methylmaleimide, N-al lylamine, N-(2-trimethylsilyloxy)methylamine (SEM), N-3-acetoxypropylamine, N-(1-isopropyl-4-nitro-2-oxo-3-pyroline-3-yl)amine, quaternary ammonium salts, N-benzylamine, N-di(4-methoxyphenyl)methylamine, N-5-dibenzo suberylamine, N-triphenylmethyamine (Tr), N-(4-methoxyphenyl)diphenylmethylamine (MMTr), N-9-phenylfluorenylamine (PhF), N-2,7-dichloro-9-fluorenylethylamine, N-ferrocenylethylamine (Fcm), N-2-picolinoyl N'-oxide, N-1,1-dimethyliothiylamine, N-benzylideneamine, N-methoxybenzylidene amine, N-diphenylmethylethylamine, N-[2-(pyridyl)iminemethyl]ethylenelamine, N-4,4'-dimethylaminomethylethyleneamine, N,N-isopropylidenediamine, N-nitrobenzylethylamine, N-cyclhexylideneamine, N-5-cyclohexylideneamine, N-(5-chloro-2-hydroxyphenyl)methylethylamine, N-cyclohexylideneamine, N-(5,5-dimethyl-3-oxo-1-cyclohexenyl) amine, N-borane derivative, N-diphenylborinic acid derivative, N-[phenyl(pentacarbonylchromium- or tungsten) carbonyl]amine, N-copper chelate, N-zinc chelate, N-nitroamine, N-nitrosamine, amine N-oxide, diphenylphosphin amide (Dpp), dimethyliophosphinamide (Mpt), diphenylasph phosphinamide (Ppt), dialkyl phosphorodates, dibenzyl phosphorodiate, diphenylphosphoridate, benzzenesulfonylamine, o-nitrobenzenesulfonylamine (Nps), 2,4-dinitrobenzenesulfonylamine, pentachlorobenzencesulfonylamine, 2-nitro-4-methoxybenzenesulfonylamine, triphenylmethanesulfonylamine, 3-nitropyridinesulfonylamine (Nps), toluenesulfonylamine (Ts), benzenesulfonylamine, 2,3,6-trim ethyl-4-methoxybenzenesulfonylamine (Mtr), 2,4,6-trimes othoxybenzenesulfonylamine (Mtb), 2,6-dimethyl-4-meth oxybenzenesulfonylamine (Pme), 2,3,5,6-tetramethyl-4- methoxybenzenesulfonylamine (Mte), 4-methoxybenzenesulfonylamine (Mbs), 2,4,6-trimethylbenzenesulfonylamine (Mts), 2,6-dimethoxy-4-methoxybenzenesulfonylamine (Mds), 2,2,5,7,8-pentamethylchroman-6-sulfonamide (Pme), methanesulfonylamine (Ms), β-trimethylsilylethanesulfonylamine (SES), antracenesulfonylamine, 4-(4'-8'-dimethoxynaphthyl)benzenesulfonylamine (DNMBS), benzenesulfonylamine, trifluoromethylsulfonylamine, and phenacylsulfonylamine. Exemplary protecting groups are described in Protecting Groups in Organic Synthesis, T.
suitable monovalent substituents on 

Suitable divalent substituents on a saturated carbon atom of R include: —O and —S.

Suitable divalent substituents on a saturated carbon atom of an "optionally substituted" group include the following: —O, —S, —NRR', or —NHR'. Suitable divalent substituents are limited to those that result in the formation of stable or chemically feasible compounds. The term "stable", as used herein, refers to compounds that are not substantially altered when subjected to conditions for their production, detection, and, in certain embodiments, their recovery, purification, and use for one or more of the purposes disclosed herein.

In some chemical structures herein, substituents are shown attached to a bond that crosses a bond in a ring of the depicted molecule. This convention indicates that one or more of the substituents may be replaced by the ring at any available position (usually in place of a hydrogen atom of the parent structure). In cases where an atom of a ring so substituted has two substitutable positions, two groups may be present on the same ring atom. Unless otherwise indicated, when more than one substituent is present, each is defined independently of the others, and each may have a different structure. In cases where the substituent shown crossing a bond of the ring is R, this has the same meaning as if the ring were said to be "optionally described" as described in the preceding paragraph.

Suitable monovalent substituents on a substitutable carbon atom of an "optionally substituted" group are independently halogen: —CH₃, —CH₂OR, —O—CH₂OR', or —CH₂OR, or sulfur, or, notwithstanding the definition above, two independent occurrences of R, taken together with intervening
atom(s) form an unsubstituted 3-12-membered saturated, partially unsaturated, or aryl mono- or bicyclic ring having 0-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur. A substitutable nitrogen may be substituted with three R³ substituents to provide a charged ammonium moiety —N⁺(R³)₃, wherein the ammonium moiety is further complexed with a suitable counterion.

Suitable substituents on the aliphatic group of R¹ are independently halogen, —R³, —(haloR³), —OH, —OR³, —O(haloR³), —CN, —C(O)OH, —C(O)OR³, —NH₂, —NR³, —NR³₂, or —NO₂, wherein each R³ is unsubstituted or where preceded by “halo” is substituted only with one or more halogens, and is independently C₁₋₆ aliphatic, —CH₂Ph, —O(CH₂)ₐPh, or a 5-6-membered saturated, partially unsaturated, or aryl ring having 0-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

As used herein, the term “catalyst” refers to a substance the presence of which increases the rate and/or extent of a chemical reaction, while not being consumed or undergoing a permanent chemical change itself.

As used herein, the term “multidentate” refers to ligands having multiple sites capable of coordinating to a single metal center.

As used herein, the term “activating moiety” refers to a moiety comprising one or more activating functional groups. In certain embodiments, an activating moiety improves the catalytic activity of a metal complex. In some embodiments, such improved catalytic activity is characterized by higher conversion of starting materials compared to a metal complex lacking an activating moiety. In some embodiments, such improved catalytic activity is characterized by higher rate of conversion of starting materials compared to a metal complex lacking an activating moiety. In some embodiments, such improved catalytic activity is characterized by higher yield of product compared to a metal complex lacking an activating moiety.

DETAILED DESCRIPTION OF CERTAIN EMBODIMENTS

The present invention provides, among other things, unimolecular metal complexes for the copolymerization of carbon dioxide and epoxides and methods of using the same. In certain embodiments, provided metal complexes contain a metal-ligand moiety tethered to one or more activating moieties. In some embodiments, an activating moiety comprises a linker and one or more activating functional groups. In some embodiments, provided metal complexes act as polymerization catalysts. In certain embodiments, at least one activating functional group present on the tethered moiety can act as a polymerization co-catalyst and thereby increase the rate of the copolymerization.

In certain embodiments, provided metal complexes include a metal atom coordinated to a multidentate ligand and at least one activating moiety tethered to the multidentate ligand. In certain embodiments, provided metal complexes have the structure:

wherein:

M is a metal atom;

comprises a multidentate ligand;

represents one or more activating moieties attached to the multidentate ligand, where is a linker moiety covalently coupled to the ligand, each Z is an activating functional group; and m is an integer from 1 to 4 representing the number of Z groups present on an individual linker moiety.

In certain embodiments, provided metal complexes include a metal atom coordinated to a multidentate ligand and at least one activating moiety tethered to the multidentate ligand. In some embodiments, there are 1 to 10 activating moieties tethered to the multidentate ligand. In certain embodiments, there are 1 to 8 such activating moieties tethered to the multidentate ligand. In certain embodiments, there are 1 to 4 such activating moieties tethered to the multidentate ligand.

I. Activating Functional Groups

In some embodiments, an activating functional group is selected from the group consisting of neutral nitrogen-containing functional groups, cationic moieties, phosphorus-containing functional groups, and combinations of two or more of these.

I.a. Neutral Nitrogen-Containing Activating Groups

In some embodiments, one or more tethered activating functional groups on provided metal complexes are neutral nitrogen-containing moieties. In some embodiments, such moieties include one or more of the structures in Table Z-1:
wherein:

each occurrence of $R^1$, and $R^2$ is independently hydrogen or an optionally substituted radical selected from the group consisting of $C_{1-20}$ aliphatic; $C_{1-20}$ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein an $R^6$ group can be taken with an $R^1$ or $R^2$ group to form one or more optionally substituted rings;

each occurrence of $R^2$ is independently hydrogen, a hydroxyl protecting group, or an optionally substituted radical selected from the group consisting of $C_{1-20}$ aliphatic; $C_{1-20}$ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein two or more $R^1$ and $R^2$ groups can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more additional heteroatoms;

each occurrence of $R^5$ is independently hydrogen or an optionally substituted radical selected from the group consisting of $C_{1-20}$ aliphatic; $C_{1-20}$ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein two or more $R^1$ and $R^2$ groups can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more additional heteroatoms;

where $R^1$ and $R^2$ are as defined above.

In certain embodiments, $R^1$ and $R^2$ are both hydrogen. In some embodiments, only one of $R^1$ and $R^2$ is hydrogen. In certain embodiments, $R^1$ and $R^2$ are each independently an optionally substituted radical selected from the group consisting of $C_{1-20}$ aliphatic; $C_{1-20}$ heteroaliphatic; 5- to 14-membered heteroaryl, phenyl, 8- to 10-membered aryl and 3- to 7-membered heterocyclic. In certain embodiments, $R^1$ and $R^2$ are each independently an optionally substituted radical selected from the group consisting of phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein two or more $R^1$ and $R^2$ groups can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more additional heteroatoms;

In certain embodiments, $R^1$ and $R^2$ are each independently an optionally substituted radical selected from the group consisting of $C_{1-12}$ aliphatic and $C_{1-12}$ heteroaliphatic. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted $C_{1-20}$ aliphatic. In some embodiments, $R^1$ and $R^2$ are each independently optionally substituted $C_{1-12}$ aliphatic. In some embodiments, $R^1$ and $R^2$ are each independently optionally substituted $C_{1-6}$ aliphatic. In
some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted \( C_{1-10} \) heteroaromatic. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted \( C_{1-13} \) heteroaromatic. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted phenyl. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted 8- to 10-membered aryl. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted phenyl group. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted 5- to 10-membered heteroaryl group. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted 3- to 7-membered heterocyclic.

In certain embodiments, \( R^1 \) and \( R^2 \) are each independently hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, optionally substituted phenyl, or optionally substituted benzyl. In certain embodiments, \( R^1 \) and \( R^2 \) are both methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, phenyl or benzyl. In some embodiments, \( R^1 \) and \( R^2 \) are each butyl. In some embodiments, \( R^1 \) and \( R^2 \) are each isopropyl. In some embodiments, \( R^1 \) and \( R^2 \) are perfluoro. In some embodiments, \( R^1 \) and \( R^2 \) are \( \text{CF}_2\text{CF}_2 \). In some embodiments, \( R^1 \) and \( R^2 \) are each phenyl. In some embodiments, \( R^1 \) and \( R^2 \) are each benzyl.

In some embodiments, \( R^1 \) and \( R^2 \) are taken together with intervening atoms to form one or more optionally substituted rings. In certain embodiments, \( R^1 \) and \( R^2 \) are taken together to form a ring fragment selected from the group consisting of: \(-\text{C}(\text{R}^1)^2-\), \(-\text{C}(\text{R}^1)^2\text{C}(\text{R}^2)^2-\), \(-\text{C}(\text{R}^1)^2\text{C}(\text{R}^2)^2\text{C}(\text{R}^3)^2-\), \(-\text{C}(\text{R}^1)^2\text{O}(\text{R}^2)^2-\), and \(-\text{C}(\text{R}^1)^2\text{N}(\text{R}^2)^2-\). In certain embodiments, \( R^1 \) and \( R^2 \) are taken together to form a ring fragment selected from the group consisting of: \(-\text{CH}_2-\), \(-\text{CH}_2\text{CH}_2-\), \(-\text{CH}_2\text{CH}_2\text{CH}_2-\), \(-\text{CH}_2\text{OCH}_2-\), and \(-\text{CH}_2\text{NHCH}_2-\). In some embodiments, \( R^1 \) and \( R^2 \) are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle.

In specific embodiments, an N-linked amine activating functional group is selected from the group consisting of:
In some embodiments, one or more activating functional groups is an N-linked hydroxyl amine derivative:

![Chemical Structure](image)

wherein $R^1$ and $R^2$ are as defined above.

In certain embodiments, $R^1$ is hydrogen. In some embodiments, $R^1$ is an optionally substituted radical selected from the group consisting of $C_{1-12}$ aliphatic, phenyl, 8- to 10-membered aryl, and 3- to 7-membered heterocyclic. In certain embodiments, $R^1$ is a $C_{1-12}$ aliphatic. In certain embodiments, $R^1$ is a $C_{1-12}$ aliphatic. In some embodiments, $R^2$ is an optionally substituted 8- to 10-membered aryl group. In certain embodiments, $R^2$ is an optionally substituted phenyl. In some embodiments, $R^2$ is methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, phenyl or benzyl.

In certain embodiments, $R^2$ is hydrogen. In some embodiments, $R^2$ is an optionally substituted radical selected from the group consisting of $C_{1-20}$ aliphatic; $C_{1-20}$ heteroaliphatic; 5- to 14-membered heteroaryl, phenyl, 8- to 10-membered aryl and 3- to 7-membered heterocyclic. In certain embodiments, $R^2$ is an optionally substituted radical selected from the group consisting of phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

In some embodiments, an activating functional group in a provided metal complex is an amidine. In certain embodiments, such amidine activating functional groups are selected from:

![Chemical Structures](image)
where each occurrence of R¹, R² and R³ are as defined above.

In certain embodiments, R¹ and R² are hydrogen. In some embodiments, only one of R¹ and R² is hydrogen. In certain embodiments, each R¹ and R² is independently an optionally substituted 5- to 14-membered heterocyclic or aromatic polycyclic carbocycle: a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle: a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle: a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or a 2- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

In certain embodiments, R¹ and R² are each independently an optionally substituted radical selected from the group consisting of C₁₋₁₀ aliphatic; C₁₋₁₀ heteroaliphatic, 5- to 14-membered heteroaryl, phenyl, 8- to 10-membered aryl and 3- to 7-membered heterocyclic. In some embodiments, each R¹ and R² is independently an optionally substituted radical selected from the group consisting of phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle: a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle: a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

In certain embodiments, N-linked amidine groups are selected from the group consisting of:—C(R¹)₂—, —C(R¹)₂C(O)R¹—, —C(R¹)₂C(=O)₂C(R¹)₂—, —C(R¹)₂OC(R¹)₂—, and —C(R¹)₂NR²(CO)₂R¹—. In certain embodiments, R¹ and R² are taken together to form a ring fragment selected from the group consisting of:—CH₂—, —CH₂CH₂—, —CH₂CH₂CH₂—, —CH₂OCH₂—, and —CH₂NR²CH₂—. In some embodiments, R¹ and R² are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle.

In certain embodiments, R³ is H. In certain embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted 6- to 14-membered aryl. In certain embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted phenyl.

In some embodiments, one or more R¹ or R² groups are taken together with R³ and intervening atoms to form an optionally substituted ring. In certain embodiments, R¹ and R² are taken together to form an optionally substituted 5- to 6-membered ring. In some embodiments, R² and R³ are taken together to form an optionally substituted 5- to 6-membered ring optionally containing one or more additional heteroatoms. In some embodiments, R¹, R² and R³ are taken together to form an optionally substituted fused ring system. In some embodiments such rings formed by combinations of any of R¹, R² and R³ are partially unsaturated or aromatic.

In certain embodiments, an activating functional group is an N-linked amidine:

In certain embodiments, N-linked amidine groups are selected from the group consisting of:—C(R¹)₂—, —C(R¹)₂C(O)R¹—, —C(R¹)₂C(=O)₂C(R¹)₂—, —C(R¹)₂OC(R¹)₂—, and —C(R¹)₂NR²(CO)₂R¹—. In certain embodiments, R¹ and R² are taken together to form a ring fragment selected from the group consisting of:—CH₂—, —CH₂CH₂—, —CH₂CH₂CH₂—, —CH₂OCH₂—, and —CH₂NR²CH₂—. In some embodiments, R¹ and R² are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle.

In certain embodiments, R³ is H. In certain embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted 6- to 14-membered aryl. In certain embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted phenyl.

In some embodiments, one or more R¹ or R² groups are taken together with R³ and intervening atoms to form an optionally substituted ring. In certain embodiments, R¹ and R² are taken together to form an optionally substituted 5- to 6-membered ring. In some embodiments, R² and R³ are taken together to form an optionally substituted 5- to 6-membered ring optionally containing one or more additional heteroatoms. In some embodiments, R¹, R² and R³ are taken together to form an optionally substituted fused ring system. In some embodiments such rings formed by combinations of any of R¹, R² and R³ are partially unsaturated or aromatic.

In certain embodiments, an activating functional group is an N-linked amidine:

In certain embodiments, N-linked amidine groups are selected from the group consisting of:—C(R¹)₂—, —C(R¹)₂C(O)R¹—, —C(R¹)₂C(=O)₂C(R¹)₂—, —C(R¹)₂OC(R¹)₂—, and —C(R¹)₂NR²(CO)₂R¹—. In certain embodiments, R¹ and R² are taken together to form a ring fragment selected from the group consisting of:—CH₂—, —CH₂CH₂—, —CH₂CH₂CH₂—, —CH₂OCH₂—, and —CH₂NR²CH₂—. In some embodiments, R¹ and R² are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle.

In certain embodiments, R³ is H. In certain embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted 6- to 14-membered aryl. In certain embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted C₁₋₁₀ aliphatic. In some embodiments, R³ is optionally substituted phenyl.
In certain embodiments, activating functional groups are amidine moieties linked through the imine nitrogen:

\[
R^1, R^2, R^3, \quad \text{and} \quad R^4.
\]

In certain embodiments, imine-linked amidine activating functional groups are selected from the group consisting of:

\[
\begin{align*}
&\text{In certain embodiments, carbon-linked amidine activating groups are selected from the group consisting of:}
\end{align*}
\]
In some embodiments, one or more activating functional groups is a carbamate. In certain embodiments, a carbamate is N-linked:

where $R^1$ and $R^2$ are as defined above. In some embodiments, a carbamate is O-linked:

where $R^1$ and $R^2$ are as defined above.

In certain embodiments, $R^1$ and $R^2$ are both hydrogen. In some embodiments, only one of $R^1$ and $R^2$ is hydrogen. In certain embodiments, $R^1$ and $R^2$ are each independently an optionally substituted radical selected from the group consisting of C$_{1-20}$ aliphatic; C$_{1-20}$ heteroaliphatic; 5- to 14-membered heteroaryl, phenyl, 8- to 10-membered aryl and 3- to 7-membered heterocyclic. In some embodiments, $R^1$ and $R^2$ are each independently an optionally substituted radical selected from the group consisting of benzyl, a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

In certain embodiments, $R^1$ and $R^2$ are each independently an optionally substituted radical selected from the group consisting of C$_{1-12}$ aliphatic and C$_{1-12}$ heteroaliphatic. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted C$_{1-20}$ aliphatic. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted C$_{1-12}$ aliphatic. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted C$_{1-12}$ aliphatic. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted C$_{1-12}$ aliphatic. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted 8- to 10-membered aryl. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted phenyl group. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted 5- to 10-membered heteroaryl group. In some embodiments, each occurrence of $R^1$ and $R^2$ is independently an optionally substituted C$_{1-12}$ aliphatic.

In certain embodiments, $R^1$ and $R^2$ are each independently hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, optionally substituted phenyl, or optionally substituted benzyl. In certain embodiments, $R^1$ and $R^2$ are both methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, phenyl or benzyl. In some embodiments, $R^1$ and $R^2$ are each butyl. In some embodiments, $R^1$ and $R^2$ are each isopropyl. In some embodiments, $R^1$ and $R^2$ are perfluoro. In some embodiments, $R^1$ and $R^2$ are —CF$_2$CF$_2$. In some embodiments, $R^1$ and $R^2$ are each phenyl. In some embodiments, $R^1$ and $R^2$ are each benzyl.

In some embodiments, $R^1$ and $R^2$ are taken together with intervening atoms to form one or more optionally substituted rings. In certain embodiments, $R^1$ and $R^2$ are taken together to form a ring fragment selected from the group consisting of: —C($R^1$)$_2$, —C($R^1$)$_3$C($R^1$)$_2$, —C($R^1$)$_4$C($R^1$)$_3$C($R^1$)$_2$.
——C(R')OC(R')——, and —C(R')₂NR'C(R')₂——. In certain embodiments, R' and R₂ are taken together to form a ring fragment selected from the group consisting of: —CH₂——, —CH₃——, —CH₂CH₂——, —CH₂CH₃——, and —CH₃NRCH₂——. In some embodiments, R₁ and R₂ are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle. In some embodiments, R₂ is selected from the group consisting of: methyl, t-butyl, t-amyl, benzyloxoy, adamantanyl, allyl, 4-methoxycarbonylphosphanyl, 2-(methylsulfonyl)ethyl, 2-(4-biphenylyl)-prop-2-yl, 2-(trimethylsilyl)ethyl, 2-bromoethyl, and 9-fernylmethyl.

In some embodiments, an activating functional group is a guanidine or bis-guanidine group:

where each occurrence of R₁, R₂, R₃, R₄, R₅, R₆, R₇, and R₈ is independently hydrogen or an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heterocyclic ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur, wherein any two or more R₁, R₂, R₃, R₄, R₅, and R₆ groups can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more additional heteroatoms.

In certain embodiments, each occurrence of R₁, R₂, R₃, R₄, R₅, R₆, R₇, and R₈ is hydrogen or an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; 3- to 7-membered heterocyclic, phenyl, and 8- to 10-membered aryl. In some embodiments, each occurrence of R₁, R₂, R₃, R₄, R₅, R₆, R₇, and R₈ is hydrogen or an optionally substituted radical selected from the group consisting of: phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heterocyclic ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur, wherein any two or more R₁, R₂, R₃, R₄, R₅, R₆, R₇, and R₈ groups can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more additional heteroatoms.

In some embodiments, each occurrence of R₁, R₂, R₃, R₄, R₅, R₆, R₇, and R₈ is hydrogen or an optionally substituted radical selected from the group consisting of the following groups: C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heterocyclic ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur. In certain embodiments, each occurrence of R₁, R₂, R₃, R₄, R₅, R₆, R₇, and R₈ is independently hydrogen or an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heterocyclic ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur. In certain embodiments, each occurrence of R₁, R₂, R₃, R₄, R₅, R₆, R₇, and R₈ is independently hydrogen or an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heterocyclic ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur. In certain embodiments, each occurrence of R₁, R₂, R₃, R₄, R₅, R₆, R₇, and R₈ is independently hydrogen or an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heterocyclic ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.
optionally containing one or more additional heteroatoms. In some embodiments, \( R^1 \) and \( R^2 \) are taken together with intervening atoms to form an optionally substituted ring optionally containing one or more additional heteroatoms. In certain embodiments, \( R^1 \) and \( R^2 \) are taken together with intervening atoms to form an optionally substituted ring optionally containing one or more additional heteroatoms. In some embodiments, \( [R^2 \text{ and } R^2'] \) and \( [R^1 \text{ and } R^1'] \) are taken together with intervening atoms to form an optionally substituted ring optionally containing one or more additional heteroatoms. In some embodiments, three or more \( R^1, R^1', R^2, R^2', R^3, \) and \( R^2' \) groups are taken together with any intervening atoms to form optionally substituted rings. In certain embodiments, \( R^1 \) and \( R^2' \) groups are taken together to form an optionally substituted 5- or 6-membered ring. In some embodiments, three or more \( R^1 \) and/or \( R^2' \) groups are taken together to form an optionally substituted fused ring system.

In certain embodiments where an activating functional group is a guanidine or bis guanidine moiety, it is chosen from the group consisting of:

![Chemical structures](image)

In some embodiments, an activating functional group is a urea:

![Urea structure](image)

where \( R^1, \) and \( R^2' \) are as defined above.
In certain embodiments, both R₁ and R₂ are hydrogen. In some embodiments, R₁ and R₂ are two groups independently an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic, phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

In certain embodiments, R₁ and R₂ are each independently an optionally substituted radical selected from the group consisting of C₁₋₁₂ aliphatic and C₁₋₁₂ heteroaliphatic. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted C₁₋₂₀ aliphatic. In some embodiments, R₁ and R₂ are each independently an optionally substituted C₁₋₁₂ aliphatic. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted C₁₋₁₂ heteroaliphatic. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted 8- to 10-membered aryl. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted phenyl group. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted 5- to 10-membered heteroaryl group. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted 3- to 7-membered heterocyclic.

In certain embodiments, R₁ and R₂ are each independently hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, optionally substituted phenyl, or optionally substituted benzyl. In certain embodiments, R₁ and R₂ are both methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, phenyl or benzyl. In some embodiments, R₁ and R₂ are each butyl. In some embodiments, R₁ and R₂ are each isopropyl. In some embodiments, R₁ and R₂ are perfluoro. In some embodiments, R₁ and R₂ are —CF₃, —CF₂CF₃. In some embodiments, R₁ and R₂ are each phenyl. In some embodiments, R₁ and R₂ are each benzyl.

In some embodiments, R₁ and R₂ are each taken together with intervening atoms to form one or more optionally substituted rings. In certain embodiments, R₁ and R₂ are taken together to form a ring fragment selected from the group consisting of: —C(R₁)₃ —C(R₂)₄C(R₃)₅ —C(R₁)₃C(R₂)₄C(R₃)₅ —C(R₁)₃OC(R₂)₄ —C(R₁)₃NR(R₅)C(R₂)₄. In certain embodiments, R₁ and R₂ are taken together to form a ring fragment selected from the group consisting of: —CH₂(—C(R₁)₃ —CH₂(—C(R₂)₄ —CH₂(—C(R₃)₅ —CH₂(—C(R₂)₄C(R₃)₅ —CH₂NR(R₅)CH₂. In some embodiments, R₁ and R₂ are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle.

In certain embodiments, activating functional groups are oxime or hydrazone groups:

where R¹, R², R³, and R⁴ are as defined above.

In certain embodiments, R₁ and R₂ are both hydrogen. In some embodiments, only one of R₁ and R₂ is hydrogen. In certain embodiments, R₁ and R₂ are each independently an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic, 5- to 14-membered heteroaryl, 8- to 10-membered aryl and 3- to 7-membered heterocyclic. In certain embodiments, R₁ and R₂ are each independently an optionally substituted radical selected from the group consisting of phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

In certain embodiments, R₁ and R₂ are each independently an optionally substituted radical selected from the group consisting of C₁₋₁₂ aliphatic and C₁₋₁₂ heteroaliphatic. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted C₁₋₂₀ aliphatic. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted C₁₋₁₂ heteroaliphatic. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted 8- to 10-membered aryl. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted phenyl group. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted 5- to 10-membered heteroaryl group. In some embodiments, each occurrence of R₁ and R₂ is independently an optionally substituted 3- to 7-membered heterocyclic.
In some embodiments, \( R^1 \) and \( R^2 \) are taken together with intervening atoms to form one or more optionally substituted rings. In certain embodiments, \( R^1 \) and \( R^2 \) are taken together to form a ring fragment selected from the group consisting of:

- \( \text{C}(\text{R}^1)_{2-3} \)
- \( \text{C}(\text{R}^2)_{2-3} \)
- \( \text{C}(\text{R}^1\text{R}^2)_{2-3} \)
- \( \text{C}(\text{R}^1)_{2-3}\text{OCC}(\text{R}^2)_{2-3} \)
- \( \text{C}(\text{R}^1)_{2-3}\text{OC}(\text{R}^2)_{2-3} \)

In certain embodiments, \( R^1 \) and \( R^2 \) are taken together to form an optionally substituted ring fragment consisting of:

- \( \text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\) (acyclic aliphatic),
- \( \text{CH}_2\text{CH}_2\text{CH}_2\text{CH}_2\) (cyclic aliphatic),
- \( \text{CH}_2\text{OCH}_2\text{CH}_2\) (aliphatic),
- \( \text{CH}_2\text{NRCH}_2\) (heterocyclic),

In some embodiments, \( R^1 \) and \( R^2 \) are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated in certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle.

In certain embodiments, \( R^2 \) is \( H \). In certain embodiments, \( R^2 \) is optionally substituted \( \text{C}_{1-20} \) aliphatic, and in some embodiments \( R^2 \) is optionally substituted 6- to 14-membered aryl. In certain embodiments, \( R^2 \) is optionally substituted \( \text{C}_{1-12} \) aliphatic and in some embodiments, optionally substituted \( \text{C}_{1-6} \) aliphatic. In certain embodiments, \( R^2 \) is optionally substituted phenyl.

In some embodiments, one or more \( R^1 \) or \( R^2 \) groups are taken together with \( R^1 \) and intervening atoms to form an optionally substituted ring. In certain embodiments, \( R^1 \) and \( R^2 \) are taken together to form an optionally substituted 5- or 6-membered ring. In some embodiments, \( R^1 \) and \( R^2 \) are taken together to form an optionally substituted 5- or 6-membered ring optionally containing one or more additional heteroatoms. In some embodiments, \( R^1 \), \( R^2 \), and \( R^3 \) are taken together to form an optionally substituted fused ring system. In some embodiments such rings formed by combinations of any of \( R^1 \), \( R^2 \), and \( R^3 \) are partially unsaturated or aromatic.

In certain embodiments, \( R^1 \) is \( H \). In certain embodiments, \( R^1 \) is optionally substituted \( \text{C}_{1-20} \) aliphatic, while in some embodiments \( R^1 \) is optionally substituted 6- to 14-membered aryl. In certain embodiments, \( R^1 \) is optionally substituted \( \text{C}_{1-12} \) aliphatic or in some embodiments, optionally substituted \( \text{C}_{1-6} \) aliphatic. In certain embodiments, \( R^1 \) is optionally substituted \( \text{C}_{1-12} \) acyl or in some embodiments, optionally substituted \( \text{C}_{1-6} \) acyl. In certain embodiments, \( R^1 \) is optionally substituted phenyl. In some embodiments, \( R^1 \) is a hydroxyl protecting group. In some embodiments, \( R^1 \) is a silyl protecting group.

In some embodiments, an activating functional group is an N-oxide derivative:

```
\[
\begin{align*}
\text{O} & \quad \text{O} \\
\text{O} & \quad \text{O} \\
\end{align*}
\]
```

where \( R^1 \) and \( R^2 \) are as defined above.

In certain embodiments, \( R^1 \) and \( R^2 \) are both hydrogen. In some embodiments, only one of \( R^1 \) and \( R^2 \) is hydrogen. In certain embodiments, \( R^1 \) and \( R^2 \) are each independently an optionally substituted radical selected from the group consisting of \( \text{C}_{1-20} \) aliphatic, \( \text{C}_{1-20} \) heteroaliphatic, 5- to 14-membered heteroaryl, phenyl, or 8- to 10-membered aryl and 3- to 7-membered heterocycle.

In certain embodiments, \( R^1 \) and \( R^2 \) are each independently an optionally substituted radical selected from the group consisting of \( \text{C}_{1-12} \) aliphatic and \( \text{C}_{1-12} \) heteroaliphatic. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently...
In some embodiments, one or more tethered activating functional groups on provided metal complexes are cationic moieties. In some embodiments, such moieties include one or more of the structures in Table Z-2:

<table>
<thead>
<tr>
<th>R1</th>
<th>R2</th>
<th>R3</th>
<th>R4</th>
<th>R5</th>
<th>R6</th>
<th>R7</th>
<th>R8</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

TABLE Z-2
TABLE Z-2-continued

<table>
<thead>
<tr>
<th>R1</th>
<th>R2</th>
<th>Y'</th>
<th>R11</th>
</tr>
</thead>
<tbody>
<tr>
<td>h</td>
<td>n</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>h</td>
<td>n</td>
<td>N</td>
<td>N</td>
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<td>N</td>
</tr>
<tr>
<td>h</td>
<td>n</td>
<td>N</td>
<td>N</td>
</tr>
<tr>
<td>h</td>
<td>n</td>
<td>N</td>
<td>N</td>
</tr>
</tbody>
</table>

or a combination of two or more of these,

wherein:
- each occurrence of \( R^1 \), \( R^2 \), and \( R^3 \) is as previously defined;
- \( R^4 \) is hydrogen, hydroxyl, optionally substituted \( C_{1-20} \) aliphatic;
- each occurrence of \( R^5 \) and \( R^6 \) is independently hydrogen or an optionally substituted radical selected from the group consisting of \( C_{1-20} \) aliphatic; \( C_{1-20} \) heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein \( R^5 \) and \( R^6 \) can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more heteroatoms, and an \( R^o \) group can be taken with an \( R^1 \) or \( R^2 \) group to form one or more optionally substituted rings;
- each occurrence of \( R^7 \), \( R^8 \), and \( R^{10} \) is independently hydrogen or an optionally substituted radical selected from the group consisting of \( C_{1-20} \) aliphatic; \( C_{1-20} \) heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein any two or more \( R^7 \), \( R^8 \), and \( R^{10} \) groups can be taken together with intervening atoms to form one or more optionally substituted rings;
- each occurrence of \( R^{11} \) is independently selected from the group consisting of: halogen, \( \text{NO}_2 \), \( \text{CN} \), \( \text{SR}^3 \), \( \text{SO}^2 \) \( \text{R}^3 \), \( \text{NRC}(\text{OR})^3 \), \( \text{OC}(\text{OR})^3 \), \( \text{CO}_2 \) \( \text{R}^3 \), \( \text{NCO} \), \( \text{N}_3 \), \( \text{OR}^3 \), \( \text{OC}(\text{OR})^3 \), \( \text{N}(\text{R}^3) \), \( \text{NRC}(\text{OR})^3 \), \( \text{NRC}(\text{OR})^3 \); or an optionally substituted radical selected from the group consisting of \( C_{1-20} \) aliphatic; \( C_{1-20} \) heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated,
partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; where each occurrence of R1 is independently hydrogen or an optionally substituted C1-6 aliphatic group, and where two or more adjacent R1 groups can be taken together to form an optionally substituted saturated, partially unsaturated, or aromatic 5- to 12-membered ring containing 0 to 4 heteroatoms; X is any anion, and Ring A is an optionally substituted, 5- to 10-membered heteroaryl group.

In certain embodiments, a cationic activating functional group is a protonated amine:

\[
\begin{align*}
R^1 & \quad \text{N} \quad R^2 \\
R' & \quad \text{H} \\
R'' & \quad \text{X} \\
\end{align*}
\]

where R1 and R2 are as defined above.

In certain embodiments, R1 and R2 are both hydrogen. In some embodiments, only one of R1 and R2 is hydrogen. In certain embodiments, R1 and R2 are each independently an optionally substituted radical selected from the group consisting of C1-20 aliphatic, C1-20 heteroaliphatic, 5- to 14-membered heteroaryl, phenyl, or 8- to 10-membered aryl and 3- to 7-membered heterocyclic. In certain embodiments, R1 and R2 are each independently an optionally substituted radical selected from the group consisting of phenyl, a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

In certain embodiments, R1 and R2 are each independently an optionally substituted radical selected from the group consisting of C1-12 aliphatic and C1-12 heteroaliphatic. In some embodiments, each occurrence of R1 and R2 is independently an optionally substituted C1-20 aliphatic. In some embodiments, R1 and R2 are each independently optionally substituted C1-6 aliphatic. In some embodiments, each occurrence of R1 and R2 is independently an optionally substituted C1-20 aliphatic. In some embodiments, each occurrence of R1 and R2 is independently an optionally substituted C1-12 aliphatic. In some embodiments, each occurrence of R1 and R2 is independently an optionally substituted 8- to 10-membered aryl.
In certain embodiments, an activating functional group is a guanidinium group:

In some embodiments, each of \( R^4, R^5, R^6, R^7, \) and \( R^8 \) is hydrogen. In some embodiments, each occurrence of \( R^4, R^5, R^6, R^7, \) and \( R^8 \) is independently hydrogen or \( C_{1-20} \) aliphatic. In some embodiments, each occurrence of \( R^4, R^5, R^6, R^7, \) and \( R^8 \) is independently hydrogen or \( C_{1-12} \) aliphatic. In some embodiments, each occurrence of \( R^4, R^5, R^6, R^7, \) and \( R^8 \) is independently hydrogen or \( C_{1-20} \) heterocyclic. In some embodiments, each occurrence of \( R^4, R^5, R^6, R^7, \) and \( R^8 \) is independently hydrogen or phenyl. In some embodiments, each occurrence of \( R^4, R^5, R^6, R^7, \) and \( R^8 \) is independently hydrogen or 8- to 10-membered aryl. In some embodiments, each occurrence of \( R^4, R^5, R^6, R^7, \) and \( R^8 \) is independently hydrogen or 5- to 10-membered heterocyclic. In some embodiments, one or more of \( R^4, R^5, R^6, \) and \( R^7 \) is optionally substituted \( C_{1-12} \) aliphatic. In certain embodiments, any of \( R^4 \) and \( R^5 \), \( R^6 \) and \( R^8 \), \( R^7 \) and \( R^8 \), \( R^4 \) and \( R^7 \), and \( R^5 \) and \( R^7 \) can be taken together with intervening atoms to form one or more optionally substituted rings. In some embodiments, \( R^4 \) and \( R^6 \) and \( R^7 \) and \( R^8 \) are taken together to form rings.

It will be appreciated that when a guanidinium cation is depicted as

all such resonance forms are contemplated and encompassed by the present disclosure. For example, such groups can also be depicted as

or
In specific embodiments, a guanidinium activating functional group is selected from the group consisting of:

In some embodiments, each occurrence of $R^8$, $R^9$, and $R^{10}$ is independently optionally substituted C$_{1,20}$ aliphatic. In some embodiments, each occurrence of $R^8$, $R^9$, and $R^{10}$ is independently hydrogen or optionally substituted C$_{1,20}$ heteroaliphatic. In some embodiments, each occurrence of $R^8$, $R^9$, and $R^{10}$ is independently hydrogen or optionally substituted phenyl. In some embodiments, each occurrence of $R^8$, $R^9$, and $R^{10}$ is independently hydrogen or optionally substituted 8- to 10-membered heterocyclic. In some embodiments, each occurrence of $R^8$, $R^9$, and $R^{10}$ is independently hydrogen or optionally substituted 5- to 10-membered heteroaryl. In some embodiments, each occurrence of $R^8$, $R^9$, and $R^{10}$ is independently hydrogen or optionally substituted 3- to 7-membered heterocyclic. In some embodiments, $R^8$ and $R^9$ are taken together with intervening atoms to form one or more rings selected from the group consisting of: optionally substituted C$_3$-C$_{14}$ carbocycle, optionally substituted 3- to 14-membered heterocycle, optionally substituted C$_5$-C$_{10}$ aryl, and optionally substituted 5- to 10-membered heteroaryl.

In certain embodiments, $R^8$, $R^9$, and $R^{10}$ are each methyl. In certain embodiments, $R^8$, $R^9$, and $R^{10}$ are each phenyl.

In specific embodiments, an arsonium activating functional group is selected from the group consisting of:

In some embodiments, a nitrogen-containing heterocycle includes a quaternized nitrogen atom. In certain embodiments, a nitrogen-containing heterocycle includes an iminium moiety such as

In some embodiments, the optionally substituted nitrogen-containing heterocycle is selected from the group consisting of: pyridine, imidazole, pyrroldine, pyrazole, quinoline, thiazole, dithiazole, oxazole, triazole, pyrazolom, isoxazole, isothiazole, tetrazole, pyrazine, thiazine, and triazine.

In certain embodiments, a nitrogen-containing heterocycle is linked to a metal complex via a ring nitrogen atom. In some embodiments, a ring nitrogen to which the attachment is made is thereby quaternized, and in some embodiments, linkage to a metal complex takes the place of an N—H bond and the nitrogen atom thereby remains neutral. In certain embodi-
ments, an optionally substituted N-linked nitrogen-containing heterocycle is a pyridinium derivative. In certain embodiments, optionally substituted N-linked nitrogen-containing heterocycle is an imidazolium derivative. In certain embodiments, optionally substituted N-linked nitrogen-containing heterocycle is a thiazolium derivative. In certain embodiments, optionally substituted N-linked nitrogen-containing heterocycle is a pyridinium derivative.

In some embodiments, an activating functional group is

\[ \text{A} \]

In certain embodiments, ring A is an optionally substituted, 5- to 10-membered heteroaryl group. In some embodiments, ring A is an optionally substituted, 6-membered heteroaryl group. In some embodiments, ring A is a ring of a fused heterocycle. In some embodiments, ring A is an optionally substituted pyridyl group.

In some embodiments, \( R^{12} \) is hydrogen. In some embodiments, \( R^{12} \) is an optionally substituted \( C_{1-20} \) aliphatic group. In some embodiments, \( R^{12} \) is \( C_{1-20} \) heteroaliphatic. In some embodiments, \( R^{12} \) is optionally substituted phenyl, 8- to 10-membered aryl, 5- to 10-membered heteroaryl. In some embodiments, \( R^{12} \) is 3- to 7-membered heterocyclic. In some embodiments, \( R^{12} \) is an optionally substituted \( C_{1-12} \) aliphatic group. In some embodiments, \( R^{12} \) is neopentyl. In some embodiments, \( R^{12} \) is oxide or hydroxyl.

In some embodiments, when \( Z \) is

\[ \text{A} \]

ring A is other than an imidazole, an oxazole, or a thiazole.

In specific embodiments, a nitrogen-containing heterocycle activating functional group is selected from the group consisting of:
In some embodiments, an activating functional group is

\[
\begin{align*}
N & \quad R^1 \quad R^2 \\
N & \quad R^1 \quad R^2 \\
N & \quad R^1 \quad R^2
\end{align*}
\]

where \(R^1, R^2,\) and \(R^3\) are as defined above.

In certain embodiments, \(R^1, R^2,\) and \(R^3\) are each hydrogen. In some embodiments, only one of \(R^1\) and \(R^2\) is hydrogen. In certain embodiments, \(R^1\) and \(R^2\) are each independently an optionally substituted radical selected from the group consisting of \(\text{C}_{1-20} \text{ aliphatic; C}_{1-12} \text{ heteroaliphatic; 5- to 14-membered heteroaryl, phenyl, 8- to 10-membered aryl and 3- to 7-membered heterocyclic. In certain embodiments, } R^1 \text{ and } R^2 \text{ are each independently an optionally substituted radical selected from the group consisting of phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.}

In certain embodiments, \(R^1, R^2,\) and \(R^3\) are each independently an optionally substituted radical selected from the group consisting of \(\text{C}_{1-12} \text{ aliphatic; C}_{1-12} \text{ heteroaliphatic. In some embodiments, each occurrence of } R^1 \text{ and } R^2 \text{ is independently an optionally substituted } \text{C}_{1-20} \text{ aliphatic. In some embodiments, each occurrence of } R^1 \text{ and } R^2 \text{ is independently an optionally substituted phenyl group. In some embodiments, each occurrence of } R^1 \text{ and } R^2 \text{ is independently an optionally substituted 5- to 10-membered heteroaryl group. In some embodiments, each occurrence of } R^1 \text{ and } R^2 \text{ is independently an optionally substituted 3- to 7-membered heterocyclic. In certain embodiments, } R^1 \text{ and } R^2 \text{ are each independently hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, optionally substituted phenyl, or optionally substituted benzyl. In certain embodiments, } R^1 \text{ and } R^2 \text{ are both methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, phenyl or benzyl. In some embodiments, } R^1 \text{ and } R^2 \text{ are each butyl. In some embodiments, } R^1 \text{ and } R^2 \text{ are each isopropyl. In some embodiments, } R^1 \text{ and } R^2 \text{ are perfluoro. In some embodiments, } R^1 \text{ and } R^2 \text{ are each phenyl. In some embodiments, } R^1 \text{ and } R^2 \text{ are each benzyl. In some embodiments, } R^1 \text{ and } R^2 \text{ are taken together with intervening atoms to form one or more optionally substituted rings. In certain embodiments, } R^1 \text{ and } R^2 \text{ are taken together to form a ring fragment selected from the group consisting of: } -C(R^1)\text{CF}_2-C(R^2)\text{CF}_2-; -C(R^1)\text{CF}_2-C(R^2)\text{CF}_2-; -C(R^1)\text{OC}(R^2)\text{CF}_2-; -C(R^1)\text{NR}(R^2)\text{CF}_2-; -CH_2\text{CF}_2-; -CH_2\text{CH}_2\text{CF}_2-; -CH_2\text{CH}_2\text{CH}_2\text{CF}_2-; -CH_2\text{OCH}_2-; \text{ and } -CH_2\text{NR}(R^1)\text{CH}_2-\text{NR}(R^2)\text{CH}_2-.\text{ In some embodiments, } R^1 \text{ and } R^2 \text{ are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle. In certain embodiments, } R^3 \text{ is H. In certain embodiments, } R^3 \text{ is optionally substituted } \text{C}_{1-20} \text{ aliphatic, and in some embodiments } R^3 \text{ is optionally substituted 6- to 14-membered aryl. In certain embodiments, } R^3 \text{ is optionally substituted } \text{C}_{1-20} \text{ aliphatic and in some embodiments, optionally substituted } \text{C}_{1-6} \text{ aliphatic. In certain embodiments, } R^2 \text{ is methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, phenyl or benzyl. In some embodiments, } R^2 \text{ is perfluoro. In some embodiments, } R^2 \text{ is } -\text{CF}_2\text{CF}_2-. In some embodiments, } R^2 \text{ is optionally substituted phenyl. In some embodiments, one or more } R^1 \text{ or } R^2 \text{ groups are taken together with } R^3 \text{ and intervening atoms to form an optionally substituted ring. In certain embodiments, } R^1 \text{ and } R^2 \text{ are taken together to form an optionally substituted 5- or 6-membered ring. In some embodiments, } R^2 \text{ and } R^3 \text{ are taken together to form an optionally substituted 5- or 6-membered ring optionally containing one or more additional heteroatoms. In some embodiments, } R^1, R^2 \text{ and } R^3 \text{ are taken together to form an optionally substituted fused ring system. In some embodiments such rings formed by combinations of any of } R^1, R^2 \text{ and } R^3 \text{ are partially unsaturated or aromatic.}

In some embodiments, an activating functional group is

\[
\begin{align*}
N & \quad R^1 \quad R^2 \\
N & \quad R^1 \quad R^2 \\
N & \quad R^1 \quad R^2
\end{align*}
\]

where \(R^1\) and \(R^2\) are as defined above.
In certain embodiments, R₃ and R⁴ are each independently an optionally substituted group selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; and 8-10-membered aryl. In some embodiments, R³ and R⁴ are each independently an optionally substituted 4-7-membered heterocyclic. In some embodiments, R³ and R⁴ can be taken together with intervening atoms to form one or more rings selected from the group consisting of: optionally substituted C₃₋₇ carbocycle, optionally substituted C₅₋₇ heterocycle, optionally substituted C₈₋₁₀ ary1, and optionally substituted 5- to 10-membered heteroaryl. In some embodiments, each occurrence of R³ and R⁴ is independently perfluoro. In some embodiments, each occurrence of R³ and R⁴ is independently —CF₃CF₂.

In some embodiments, an activating functional group is

\[ \text{R', R, R, and R are as defined above.} \]

In certain embodiments, R¹, R₂, and R⁵ are each independently an optionally substituted group selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; and 8-10-membered aryl. In certain embodiments, R¹, R₂, and R⁵ are each independently an optionally substituted 4-7-membered heterocyclic. In some embodiments, R¹ and R₂ can be taken together with intervening atoms to form one or more rings selected from the group consisting of: optionally substituted C₃₋₇ carbocycle, optionally substituted C₅₋₇ heterocycle, optionally substituted C₈₋₁₀ ary1, and optionally substituted 5- to 10-membered heteroaryl. In certain embodiments, R¹, R₂, and R⁵ are each independently an optionally substituted radical selected from the group consisting of C₁₋₄ aliphatic; C₁₋₄ heteroaliphatic; phenyl; and 8-10-membered aryl. In some embodiments, R¹ and R₂ are each independently an optionally substituted 4-7-membered heterocyclic. In some embodiments, R¹ and R₂ can be taken together with intervening atoms to form one or more rings selected from the group consisting of: optionally substituted C₃₋₇ carbocycle, optionally substituted C₅₋₇ heterocycle, optionally substituted C₈₋₁₀ ary1, and optionally substituted 5- to 10-membered heteroaryl. In some embodiments, R¹ and R₂ are each independently selected from the group consisting of methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, or benzy1. In some embodiments, each occurrence of R¹ and R₂ is independently perfluoro. In some embodiments, each occurrence of R¹ and R₂ is independently —CF₃CF₂.

In certain embodiments, R² and R⁵ are each independently an optionally substituted group selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; and 8-10-membered aryl. In some embodiments, R¹ and R² are each independently an optionally substituted 4-7-membered heterocyclic. In some embodiments, R¹ and R² can be taken together with intervening atoms to form one or more rings selected from the group consisting of: optionally substituted C₃₋₇ carbocycle, optionally substituted C₅₋₇ heterocycle, optionally substituted C₈₋₁₀ ary1, and optionally substituted 5- to 10-membered heteroaryl. In some embodiments, R¹ and R² are each independently selected from the group consisting of methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, or benzy1. In some embodiments, each occurrence of R¹ and R² is independently perfluoro. In some embodiments, each occurrence of R¹ and R² is independently —CF₃CF₂.
sisting of C_{1-12} aliphatic and C_{1-12} heteroaliphatic. In some embodiments, each occurrence of R' and R" is independently an optionally substituted C_{1-20} aliphatic. In some embodiments, R' and R" are each independently optionally substituted C_{1-12} aliphatic. In some embodiments, R' and R" are each independently optionally substituted C_{1-6} aliphatic. In some embodiments, each occurrence of R' and R" is independently an optionally substituted C_{1-20} heteroaliphatic. In some embodiments, each occurrence of R' and R" is independently an optionally substituted C_{1-12} heteroaliphatic. In some embodiments, each occurrence of R' and R" is independently an optionally substituted C_{1-6} heteroaliphatic. In some embodiments, each occurrence of R' and R" is individually an optionally substituted phenyl or 8- to 10-membered aryl. In some embodiments, each occurrence of R' and R" is independently an optionally substituted phenyl group. In some embodiments, each occurrence of R' and R" is independently an optionally substituted 5- to 10-membered heteroaryl ring. In some embodiments, each occurrence of R' and R" is independently an optionally substituted 3- to 7-membered heterocyclic.

In certain embodiments, R', R", and R are each independently hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, optionally substituted phenyl, or optionally substituted benzyl. In certain embodiments, R' and R" are both methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, phenyl or benzyl. In some embodiments, R', R", and R" are each butyl. In some embodiments, R', R", and R" are each isopropyl. In some embodiments, R', R", and R" are perfluoro. In some embodiments, R', R", and R" are —CF₃. In some embodiments, R', R", and R" are each phenyl. In some embodiments, R', R", and R" are each benzyl.

In some embodiments, R', R", and R" are taken together with intervening atoms to form one or more optionally substituted rings. In certain embodiments, R', R", and R" are taken together to form a ring fragment selected from the group consisting of: —C(R')₂ —, —C(R')₂C(R')₂ —, —C(R')₂C(R')₂ —, —C(R')₂C(R')₂ —, —C(R')₂OC(R')₂ —, and —C(R')₂NR'C(R')₂ —. In certain embodiments, R' and R" are taken together to form a ring fragment selected from the group consisting of: —CH₂ —, —CH₂CH₂ —, —CH₂CH₂CH₂ —, —CH₂OCH₂ —, and —CH₂NR'C —. In some embodiments, R', R", and R" are each taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring is partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused poly cyclic heterocycle. In some embodiments, an activating functional group is

\[ \text{R}^1 \text{R}^2 \]

In certain embodiments, R', R", and R" are each independently an optionally substituted group selected from the group consisting of C_{1-20} aliphatic; C_{1-20} heteroaliphatic; phenyl; and 8-10-membered aryl. In certain embodiments, R', R", and R" are each independently an optionally substituted 4-7-membered heterocyclic. In some embodiments, R' and R" can be taken together with intervening atoms to form one or more rings selected from the group consisting of: optionally substituted C₅₋₁₀ carbocycle, optionally substituted C₅₋₁₀ heterocycle, optionally substituted C₅₋₁₀ aryl, and optionally substituted 5- to 10-membered heteroaryl. In certain embodiments, R', R", and R" are each independently an optionally substituted C_{1-20} aliphatic. In certain embodiments, R', R", and R" are each independently selected from the group consisting of methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, or benzyl. In certain embodiments, R', R", and R" are each independently perfluoro. In some embodiments, R', R", and R" are each independently —CF₃.

In some embodiments, an activating functional group is

\[ \text{R}^1 \text{R}^2 \]

where R' and R" are as defined above.

In certain embodiments, R', R", and R" are each hydrogen. In some embodiments, only one of R' and R" is hydrogen. In certain embodiments, R' and R" are each independently an optionally substituted radical selected from the group consisting of C_{1-20} aliphatic; C_{1-20} heteroaliphatic; phenyl; a 3- to 8-membered saturated or unsaturated monocyclic carboxylate, a 7-14 carbon saturated, partially unsaturated or aromatic monocyclic carboxylate; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.
In certain embodiments, \( R^1 \) and \( R^2 \) are each independently an optionally substituted radical selected from the group consisting of \( C_{1-12} \) aliphatic and \( C_{1-12} \) heteroaliphatic. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted \( C_{1-20} \) aliphatic. In some embodiments, \( R^1 \) and \( R^2 \) are each independently optionally substituted \( C_{1-12} \) aliphatic. In some embodiments, \( R^1 \) and \( R^2 \) are each independently optionally substituted \( C_{4,6} \) aliphatic. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted \( C_{1-12} \) heteroaliphatic. In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently an optionally substituted \( 5\text{-}10\)-membered heterocycle.

In certain embodiments, \( R' \) and \( R \) are each independently optionally Substituted group selected from the group consisting of optionally Substituted C-C carbocycle, optionally Substituted C-C heterocycle, optionally Substituted C-C aryl, and optionally Substituted 5- to 10-membered heteroaryl. In some embodiments, each occurrence of \( R' \) and \( R \) is independently an optionally substituted phenyl group. In some embodiments, each occurrence of \( R'^1 \) and \( R'^2 \) is independently an optionally substituted \( C_{1-12} \) heteroaliphatic. In some embodiments, each occurrence of \( R' \) and \( R \) is independently an optionally substituted 5- to 10-membered heteroaryl group. In some embodiments, each occurrence of \( R' \) and \( R \) is independently an optionally substituted 5- to 7-membered heterocycle.

In certain embodiments, \( R^1 \) and \( R^2 \) are each independently optionally hydrogen, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, optionally substituted phenyl, or optionally substituted benzyl. In certain embodiments, \( R^1 \) and \( R^2 \) are both alkyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, phenyl or benzyl. In some embodiments, \( R^1 \) and \( R^2 \) are each butyl. In some embodiments, \( R^1 \) and \( R^2 \) are each isopropyl. In some embodiments, \( R^1 \) and \( R^2 \) are each fluorine. In some embodiments, \( R^1 \) and \( R^2 \) are each phenyl. In some embodiments, \( R^1 \) and \( R^2 \) are each phenyl.

In some embodiments, \( R^1 \) and \( R^2 \) are taken together with intervening atoms to form one or more optionally substituted rings. In certain embodiments, \( R^1 \) and \( R^2 \) are taken together to form a ring fragment selected from the group consisting of:

\[-C(R')_2OC(R')_2-, -C(R')_2OC(R')_2-, -C(R')_2OC(R')_2-, -C(R')_2OC(R')_2-, -C(R'OOC(R')_2-, -C(R')_2OC(R')_2-,\]

In certain embodiments, \( R^1 \) and \( R^2 \) are taken together to form a ring fragment selected from the group consisting of:

\[-CH_2-, -CH_2CH_2-, -CH_2CH_2-, -CH_2CH_2-, -CH_2CH_2-, -CH_2CH_2-,\]

In some embodiments, \( R^1 \) and \( R^2 \) are taken together to form an unsaturated linker moiety optionally containing one or more additional heteroatoms. In some embodiments, the resulting nitrogen-containing ring partially unsaturated. In certain embodiments, the resulting nitrogen-containing ring comprises a fused polycyclic heterocycle.

In some embodiments, an activating functional group is

\[
\begin{align*}
R^1 & \quad \text{and} \quad R^2 \\
R^1 & \quad \text{and} \quad R^2
\end{align*}
\]

where \( R^1 \) and \( R^2 \) are as defined above.

In certain embodiments, \( R^1 \) and \( R^2 \) are each independently an optionally substituted group selected from the group consisting of \( C_{1-20} \) aliphatic; \( C_{1-20} \) heteroaliphatic; phenyl; and 8-10-membered aryl. In some embodiments, \( R^1 \) and \( R^2 \) are each independently an optionally substituted 4-7-membered heterocyclic. In some embodiments, \( R^1 \) and \( R^2 \) can be taken together with intervening atoms to form one or more rings selected from the group consisting of:

\[-C_3-C_{14} \text{ carbocycle}, \text{ optionally substituted } C_3-C_{14} \text{ heterocycle}, \text{ optionally substituted } C_3-C_{14} \text{ heteroar }
\]

In some embodiments, each occurrence of \( R^1 \) and \( R^2 \) is independently a fused polycyclic heterocycle.
In certain embodiments, a phosphorous-containing functional group is chosen from the group consisting of:

or a combination of two or more of these

wherein \( R^1 \) and \( R^2 \) are as defined above; and

each \( R^7 \), is independently hydrogen, a hydroxyl protecting group, or an optionally substituted radical selected from the group consisting of \( C_{1-20} \) alkyl, \( C_{1-20} \) aliphatic; \( C_{1-20} \) heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaromatic ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and where two \( R^7 \) groups can be taken together with intervening atoms to form an optionally substituted ring optionally containing one or more heteroatoms, and an \( R^7 \) group can be taken with an \( R^1 \) or \( R^2 \) group to an optionally substituted ring.

In some embodiments, an activating functional group is a phosphonate group:

\[
\begin{align*}
&= \text{phosphonate group} \\
&= \text{phosphonate group}
\end{align*}
\]

wherein \( R', R'', \text{and } R'' \) is as defined above.

In specific embodiments, a phosphonate activating functional group is selected from the group consisting of:

\[
\begin{align*}
\text{wherein } R', R'', \text{and } R''' \text{ are as defined above.}
\end{align*}
\]

In some embodiments, an activating functional group is a phosphonic diamide group:

\[
\begin{align*}
&= \text{phosphonic diamide group} \\
&= \text{phosphonic diamide group}
\end{align*}
\]

wherein \( R', R'', \text{and } R''' \) are as defined above. In certain embodiments, each \( R' \) and \( R'' \) group in a phosphonic diamide is methyl.

In some embodiments, an activating functional group is a phosphine group:

\[
\begin{align*}
&= \text{phosphine group} \\
&= \text{phosphine group}
\end{align*}
\]

In specific embodiments, a phosphine activating functional group is selected from the group consisting of:

II. Linker Moieties

As described above, each activating moiety \((Z)_m\) comprises a linker \(-\) coupled to at least one activating functional group \(Z\) as described above, with \( m \) denoting the number of activating functional groups present on a single linker moiety.
As noted above there may be one or more activating moiety (Z), tethered to a given metal complex, similarly, each activating moiety itself may contain more than one activating functional group Z. In certain embodiments, each activating moiety contains only one activating functional group (i.e. m=1). In some embodiments, each activating moiety contains more than one activating functional groups (i.e. m>1). In certain embodiments, an activating moiety contains two activating functional groups (i.e. m=2). In certain embodiments, an activating moiety contains three activating functional groups (i.e. m=3). In certain embodiments, an activating moiety contains four activating functional groups (i.e. m=4). In certain embodiments where more than one activating functional group is present on an activating moiety, two or more of the activating functional groups are different.

In certain embodiments, each linker moiety contains 1-30 atoms including at least one carbon atom, and optionally one or more atoms selected from the group consisting of N, O, S, Si, B, and P. In certain embodiments, the linker is an optionally substituted C₃₋₅ aliphatic group wherein one or more methylene units are optionally and independently replaced by —NR²—, —N(R³)C(O) —, —C(O)N(R⁴) —, —O—, —C(O) —, —OC(O) —, —S—, —SO—, —SO₂—, —C(=S)—, —C(=NR²)—, or —N—N—, where each occurrence of R² is independently —H, or an optionally substituted radical selected from the group consisting of C₁₋₅ aliphatic 3- to 7-membered heterocyclic, phenyl, and 8- to 10-membered aryl. In certain embodiments, a linker moiety is a C₄₋₁₀ aliphatic group substituted with one or more moieties selected from the group consisting of halogen, —NO₂—, —CN, —SR³—, —SO₃—, —S(O)R⁴—, —S(O)₂R⁵—, —NR³C(O)R⁶—, —OC(O)R⁷—, —CO₂R⁸—, —NCO—, —NR³—, —OR³—, —OC(O)NR³R⁸—, —N(R³)₂—, —NR³C(O)R⁶—, and —NR³C(O)OR³, where R³ is —H, —R⁹, or an optionally substituted radical selected from the group consisting of C₁₋₃ aliphatic 3- to 7-membered heterocyclic, phenyl, and 8- to 10-membered aryl.

In certain embodiments, a linker moiety is an optionally substituted C₃₋₆ aliphatic group. In certain embodiments, a linker moiety is an optionally substituted C₄₋₁₀ aliphatic group. In certain embodiments, a linker moiety is an optionally substituted C₄₋₁₀ aliphatic group. In certain embodiments, a linker moiety is an optionally substituted C₄₋₁₀ aliphatic group. In certain embodiments, a linker moiety is an optionally substituted C₄₋₁₀ aliphatic group. In certain embodiments, a linker moiety is an optionally substituted C₄₋₁₀ aliphatic group. In certain embodiments, a linker moiety is an optionally substituted C₄₋₁₀ aliphatic group. In certain embodiments, a linker moiety is an optionally substituted C₄₋₁₀ aliphatic group. In certain embodiments, a linker moiety is an optionally substituted C₄₋₁₀ aliphatic group.
III. Metal Complexes

As noted above, the present invention encompasses metal complexes that include a metal atom coordinated to a multidentate ligand and at least one activating moiety tethered to a multidentate ligand. In certain embodiments, provided metal complexes have the structure:

where * represents the site of attachment to a ligand, and each # represents a site of attachment of an activating functional group.

In some embodiments, s is 0. In some embodiments, s is 1. In some embodiments, s is 2. In some embodiments, s is 3. In some embodiments, s is 4. In some embodiments, s is 5. In some embodiments, s is 6.

In certain embodiments, t is 1. In some embodiments, t is 2. In some embodiments, t is 3. In some embodiments, t is 4.

III. Metal Atoms

In certain embodiments, M is a metal atom selected from periodic table groups 3-13, inclusive. In certain embodiments, M is a transition metal selected from periodic table groups 5-12, inclusive. In certain embodiments, M is a transition metal selected from periodic table groups 4-11, inclusive. In certain embodiments, M is a transition metal selected from periodic table groups 5-10, inclusive. In certain embodiments, M is a transition metal selected from periodic table groups 7-9, inclusive. In some embodiments, M is selected from the group consisting of Cr, Mn, V, Fe, Co, Mo, W, Ru, Al, and Ni. In some embodiments, M is a metal atom selected from the group consisting of: cobalt; chromium; aluminum; titanium; ruthenium, and manganese. In some embodiments, M is cobalt. In some embodiments, M is chromium. In certain embodiments, M is aluminum.

In certain embodiments, a metal complex is a zinc, cobalt, chromium, aluminum, titanium, ruthenium, or manganese complex. In certain embodiments, a metal complex is an aluminum complex. In some embodiments, a metal complex is a chromium complex.
In some embodiments, a metal complex is a zinc complex. In certain some embodiments, a metal complex is a titanium complex. In some embodiments, a metal complex is a ruthenium complex. In certain embodiments, a metal complex is a manganese complex. In certain embodiments, a metal complex is cobalt complex. In certain embodiments where the metal complex is a cobalt complex, the cobalt metal has an oxidation state of 3+ (i.e., Co(III)). In some embodiments, the cobalt metal has an oxidation state of 2+.

III.b. Ligands

In some embodiments, a metal complex comprises a metal atom coordinated to a single tetradentate ligand and in some embodiments, the metal complex comprises a chelate containing a plurality of individual ligands. In certain embodiments, a metal complex contains two bidentate ligands. In some embodiments, a metal complex contains a tridentate ligand.

In various embodiments, tetradentate ligands suitable for metal complexes of the present invention may include, but are not limited to: salen derivatives 1, derivatives of salan ligands 2, bis-2-hydroxybenzamido derivatives 3, derivatives of the Trost ligand 4, porphyrin derivatives 5, derivatives of tetra- benzoporphyrin ligands 6, derivatives of corrole ligands 7, phthalocyaninate derivatives 8, and dibenzotetramethyltetraaza[14]anulene (tmtaa) derivatives 9 or 9'.
In some embodiments, a metal multidentate ligand coordinated with a metal complex may comprise a plurality of discrete ligands. In some embodiments, metal complexes include two bidentate ligands. In certain embodiments, such bidentate ligands may have the structure

where \( R' \) and \( R'' \) are as defined above. Metal complexes having two such ligands may adopt one of several geometries, and the present disclosure encompasses such variations.

In certain embodiments, metal complexes including two bidentate ligands may have structures selected from the group consisting of:

where each

\[ \text{represents a ligand:} \]

In certain embodiments, a tetradentate ligand is a salen ligand. In certain embodiments, a metal complex is a metallosalenate. In certain embodiments, a metal complex is a cobalt salen complex. In certain embodiments, a metal complex is a chromium salen complex. In some embodiments, a metal complex is an aluminum salen complex.

In certain embodiments, at least one activating moiety is tethered to a carbon atom of a phenyl ring of the salicylaldehyde-derived portions of a salen ligand. In certain embodiments, at least one activating moiety is tethered to a carbon atom of a porphyrin ligand. In certain embodiments, at least one activating moiety is tethered to a pyrrole-carbon atom of a porphyrin ligand. In certain embodiments, at least one activating moiety is tethered to a carbon atom forming the bridge between the pyrrole rings of a porphyrin ligand.

In certain embodiments, at least one activating moiety is tethered to one or more carbon atoms of only one phenyl ring of the salicylaldehyde-derived portions of a salen ligand, as shown in formula I:

wherein:

\[ \text{M is a metal atom;} \]
\[ \text{X is a nucleophile capable of ring opening an epoxide;} \]
\[ \text{k is an integer from 0-2 inclusive;} \]
\[ \text{R'} \text{ represents one or more substituents optionally present on the phenyl rings and each R'} \text{ is independently selected from the group consisting of: halogen, --NO, --CN; --SR', --S(O)R', --S(O)O, --NR'(O)R', --OR', --CO}, \]
\[ \text{CO}, \text{OC}, \text{OC}, \text{OR}, \text{N=}, \text{O=}, \text{OR, or} \text{or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfurs; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfurs; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfurs; or a heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfurs.} \]
sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur, where two or more adjacent R' groups can be taken together to form an optionally substituted saturated, partially unsaturated, or aromatic 5- to 12-membered ring containing 0 to 4 heteroatoms; R' is —H, or an optionally substituted radical selected from the group consisting of C₁–₆ aliphatic, 3- to 7-membered heterocyclic, phenyl, and 8- to 10-membered aryl,

represents an optionally substituted moiety linking the two nitrogen atoms of the diamine portion of the salen ligand, where

is selected from the group consisting of phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carboxylic acid; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carboxylic acid; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an optionally substituted C₂–₂₀ aliphatic group, wherein one or more methylene units are optionally and independently replaced by —NR'', —NR''—, —N(R'')C(O)−, —N(R'')N—, —OC(O)N(R'')—, —N(R'')C(O)O—, —OC(O)(O)O—, —O—, —C(=O)O—, —C(=O)O—, —C(= S)—, —C(=NR'')—, —C(=NOR'')— or —N—N—;

—(Z)ₘ represents one or more activating moieties, where “” is a covalent linker containing one or more atoms selected from the group consisting of C, O, N, S, and Si; Z is an activating functional group and m is an integer from 1 to 4 indicating the number of individual activating functional groups present in each activating moiety.

In certain embodiments, both salicylaldehyde-derived portions of a salen ligand bear one or more activating moieties:

wherein M, X, k, R',

and —(Z)ₘ are as defined above.

In some embodiments, provided metal complexes comprise a

moiety that has the structure:

wherein:

M is a metal atom,

R₁₅, R₁₆, R₁₇, R₁₈, R₁₉, R₂₀, R₂₁, and R₂₂ are independently a —(Z)ₘ group, hydrogen, halogen, —OR, —NR, —SR, —CN, —NO₂, —SO₂R, —SOR, —SO₃NR₆; —CNO, —NRSO₂R', —NCO, —N=SiR₃; or an optionally substituted radical selected from the group consisting of C₁–₂₀ aliphatic; C₁–₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carboxylic acid; a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carboxylic acid; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an optionally substituted C₂–₂₀ aliphatic group, wherein one or more methylene units are optionally and independently replaced by —NR'', —NR''—, —N(R'')C(O)−, —N(R'')N—, —OC(O)N(R'')—, —N(R'')C(O)O—, —OC(O)(O)O—, —O—, —C(=O)O—, —C(=S)—, —C(=NR'')—, —C(=NOR'')— or —N—N—;

each R is independently hydrogen, an optionally substituted radical selected from the group consisting of acyl; carboxyl; alkyl; aryl; phenyl; 8- to 10-membered aryl; C₁–₁₂ aliphatic; C₁–₁₂ heteroaliphatic; 5- to 10-membered heteroaryl; 4- to 7-membered heterocyclic; an oxygen protecting group; and a nitrogen protecting group; or:
two R on the same nitrogen atom are taken with the nitrogen to form a 3- to 7-membered heterocyclic ring;

wherein any of [R₁₋₅], [R₁₋₅ and R₁₋₅], [R₁₋₅ and R₁₋₅], and [R₁₋₅ and R₁₋₅] may optionally be taken together with the carbon atoms to which they are attached to form one or more rings which may be in turn be substituted with one or more R₁₋₅ groups; and
R^{aa} is selected from the group consisting of:

a) \[
\begin{array}{c}
\includegraphics[width=0.1\textwidth]{image1.png}
\end{array}
\]

b) \[
\begin{array}{c}
\includegraphics[width=0.1\textwidth]{image2.png}
\end{array}
\]

c) \[
\begin{array}{c}
\includegraphics[width=0.1\textwidth]{image3.png}
\end{array}
\]

d) \[
\begin{array}{c}
\includegraphics[width=0.1\textwidth]{image4.png}
\end{array}
\]

where

R^a at each occurrence is independently a \(\ldots\ Z\) group, hydrogen, halogen, \(-OR\), \(-SR\), \(-CN\), \(-NO_2\), \(-SO_2R\), \(-SOR\), \(-SO_2NR_2\); \(-CNO\), \(-NRSO_2R\), \(-NCO\), \(-N_3\), \(-SiR_3\); or an optionally substituted radical selected from the group consisting of C_{1-20} aliphatic; C_{1-20} heteroaliphatic; phenyl; a 3- to 8-membered saturated or unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur;

where:

two or more R^a groups may be taken together with the carbon atoms to which they are attached and any intervening atoms to form one or more rings;

two R^a groups are attached to the same carbon atom, they may be taken together along with the carbon atom to which they are attached to form a moiety selected from the group consisting of: a 3- to 8-membered spirocyclic ring, a carbonyl, an oxime, a hydrazine, an imine;

X is a nucleophile capable of ring opening an epoxide;

Y is a divalent linker selected from the group consisting of:

\(-NR\), \(-N(R)C(O)\), \(-C(O)NR\), \(-O\), \(-C(O)\), \(-OC(O)\), \(-C(O)O\), \(-S\), \(-SO\), \(-SO_2\), \(-C(=S)\), \(-C(=NR)\), or \(-N\equiv N\); a polyether; a C_3 to C_8 substituted or unsubstituted carbocycle; and a C_1 to C_8 substituted or unsubstituted heterocycle;
m' is 0 or an integer from 1 to 4, inclusive;
q is 0 or an integer from 1 to 4, inclusive; and
x is 0, 1, or 2.

In some embodiments, at least one of [R^{aa} and R^{aa'}] and [R^{aa'} and R^{aa''}] are taken together to form a ring. In some embodiments, both [R^{aa} and R^{aa'}] and [R^{aa'} and R^{aa''}] are taken together to form rings. In some embodiments, the rings formed by [R^{aa} and R^{aa'}] and [R^{aa'} and R^{aa''}] are substituted phenyl rings.

In certain embodiments, one or more of R^{1a}, R^{1a'}, R^{2a}, R^{2a'}, R^{3a}, and R^{3a'} are independently a \(\ldots\ Z\) group.

In certain embodiments of provided metal complexes, a moiety has a structure selected from the group consisting of:

[diagram of metal complex structure]
partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur;

wherein [R⁻¹ and R⁺], [R⁻² and R⁺] and any two adjacent R⁻³, R⁻⁴, R⁻⁵, R⁻⁶, R⁻⁷, and R⁻⁸ groups can be taken together with intervening atoms to form one or more optionally substituted rings;

n is 0 or an integer from 1 to 8, inclusive; and

p is 0 or an integer from 1 to 4, inclusive.

In some embodiments, M is Co.

In some embodiments, R⁻¹, R⁻², R⁻³, R⁻⁴, R⁻⁵, R⁻⁶, and R⁻⁷ are each —H. In some embodiments, R⁻¹, R⁻², R⁻³, and R⁻⁷ are each optionally substituted C₁-C₁₂ aliphatic. In some embodiments, R⁻¹, R⁻², R⁻³, R⁻⁶, R⁻⁷, and R⁻⁸ are each independently selected from the group consisting of: —H, —SiR₃; methyl, ethyl, n-propyl, i-propyl, n-butyl, sec-butyl, t-butyl, isoamyl, t-amyl, hexyl, and trityl. In some embodiments, R⁻¹, R⁻², R⁻³, R⁻⁶, R⁻⁷, and R⁻⁸ are each —H. In some embodiments, R⁻³ is selected from the group consisting of: —H; methyl; ethyl; n-propyl; i-propyl; n-butyl; sec-butyl; t-butyl; isoamyl; t-amyl; hexyl; and trityl. In some embodiments, R⁻³ and R⁻⁸ are independently selected from the group consisting of: —H; methyl; ethyl; n-propyl; i-propyl; n-butyl; sec-butyl; t-butyl; isoamyl; t-amyl; hexyl; and trityl. In certain embodiments, one or more of R⁻³, R⁻⁶, R⁻⁷, and R⁻⁸ is a —Z group. In some embodiments, R⁻³ and R⁻⁸ are a —Z group.

In certain embodiments of metal complexes, a moiety has a structure selected from the group consisting of:
where \((Z)_n\) represents one or more independently-defined activating moieties which may be bonded to any one or more unsubstituted positions of a salicylaldehyde-derived phenyl ring.

In certain embodiments, there is an activating moiety tethered to the position ortho to a metal-bound oxygen substituent of one or both of the salicylaldehyde-derived phenyl rings of a salen ligand as in formulae IIIa and IIIb:
wherein:
M, X, k, R',

and (Z)_n are as defined above, and
R^{4a}, R^{4a'}, R^{5a}, R^{5a'}, R^{5b}, and R^{5b'} are each independently a Z group, hydrogen, halogen, —OR, —NR, —SR, —CN, —NO, —SO, —SOR, —SO, NR, —CNO, —NRSO, —NCO, —N, —SiR, or an optionally substituted radical selected from the group consisting of C_{1-20} aliphatic; C_{1-20} heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur;

wherein any two adjacent R^{4a}, R^{4a'}, R^{5a}, R^{5a'}, R^{5b}, and R^{5b'} groups can be taken together with intervening atoms to form one or more optionally substituted rings.

In certain embodiments of compounds having formulae IIIa or IIIb, R^{4a}, R^{4a'}, R^{5a}, and R^{5a'} are each hydrogen, and R^{5b}, R^{5b'} are, independently, optionally substituted C_{1-20} aliphatic.

In certain embodiments of complexes IIIa and IIIb, at least one of the phenyl rings comprising a salicylaldehyde-derived portion of a catalyst is independently selected from the group consisting of:
In certain embodiments, there is an activating moiety tethered to the position para to the phenolic oxygen of one or both of a salicylaldehyde-derived phenyl rings of the salen ligand as in structures IVa and IVb:

where M, X, k, R', R'6a, R'6s, R'6a', R'7a, R'7a', and \((Z)_m\) are as defined above.

In certain embodiments of compounds having formulae IVa or IVb, R''6a, R''6s, R''6a', and R''7a' are hydrogen, and each R''7a, R''7a' is, independently, optionally substituted C1-C20, aliphatic.

In certain embodiments of catalysts IVa and IVb, at least one of the phenyl rings comprising a salicylaldehyde-derived portion of a catalyst is independently selected from the group consisting of:
In some embodiments, there is an activating moiety tethered to the position para to the imine substituent of one or both of the salicylaldehyde-derived phenyl rings of a salen ligand as in formulae Va or Vb:

where \( M, X, k, R', R^{4a}, R^{4e}, R^{5a}, R^{5e}, R^7a, R^{7a} \), and \( (Z)_n \) are as defined above.

In certain embodiments of compounds having formulae Va or Vb, each \( R^a \) and \( R^{4a} \) is hydrogen, and each \( R^{5a}, R^{7a}, R^{7a'} \) is, independently, hydrogen or optionally substituted \( C_1-C_{20} \) aliphatic.

In certain embodiments of catalysts Va and Vb, at least one of the phenyl rings comprising a salicylaldehyde-derived portion of a catalyst is independently selected from the group consisting of:
In some embodiments, there is an activating moiety tethered to the position ortho to the imine substituent of one or both of the salicylaldehyde-derived phenyl rings of a salen ligand as in formulae VIa and VIb:

where X, k, M, R', R'^5, R'5', R'^6, R'^6', R'7, R'7',

and (Z)m are as defined above.

In certain embodiments of compounds having formulae VIa or VIb, each R'^6 and R''6 is hydrogen, and each R'^5, R'5', R'^6, R'^6', R'7, and R'7' is, independently, hydrogen or optionally substituted C1-C20 aliphatic.

In certain embodiments of catalysts VIa and VIb, at least one of the phenyl rings comprising a salicylaldehyde-derived portion of a catalyst is independently selected from the group consisting of:
In some embodiments, there are activating moieties tethered to the positions ortho and para to the phenolic oxygen of one or both of the salicylaldehyde-derived phenyl rings of a salen ligand as in formulae VIIa and VIIb:

VIIa

VIIb

where $X$, $k$, $M$, $R'$, $R^5$, $R^{5'}$, $R^7$, and $R^{7'}$ is, independently, hydrogen or optionally, hydrogen or substituted $C_1$-$C_{20}$ aliphatic.

In certain embodiments of compounds having formulae VIIIa or VIIIb, each $R^4$, $R^4'$, $R^6$, and $R^{6'}$ is hydrogen.

In some embodiments, there are activating moieties tethered to the positions ortho and para to the imine substituent of one or both of the salicylaldehyde-derived phenyl rings of a salen ligand as in formulae VIIIa and VIIIb:
In some embodiments, there is an activating moiety tethered to the imine carbon of a salen ligand as in formulae IXa and IXb:
where $M, X, k, R^{6a}, R^{6a'}, R^{5a}, R^{5a'}, R^{6b}, R^{6b'}, R^{7a}, R^{7a'}, \ldots$ are as defined above with the proviso that the atom of the activating moiety attached to the salen ligand is a carbon atom.

In certain embodiments of compounds having formulae IXa or IXb, each $R^{6a}, R^{6a'}, R^{6b},$ and $R^{6b'}$ is hydrogen, and each $R^{5a}, R^{5a'}, R^{5b},$ and $R^{5b'}$ is, independently, hydrogen or optionally substituted $C_1-C_{20}$ aliphatic.

In certain embodiments of the present invention, metal complexes of structures IXa or IXb above, at least one of the phenyl rings comprising a salicylaldehyde-derived portion of a catalyst is independently selected from the group consisting of:
As shown above, the two phenyl rings derived from salicylaldehyde in the core salen structures need not be the same. Though not explicitly shown in formulae Ia through IXb above, it is to be understood that a catalyst may have an activating moiety attached to different positions on each of the two rings, and such compounds are specifically encompassed within the scope of the present invention. Furthermore, activating moieties can be present on multiple parts of the ligand, for instance activating moieties can be present on the diamine bridge and on one or both phenyl rings in the same catalyst.

In certain embodiments, the salen ligand cores of catalysts Ia through IXb above are selected from the group shown below wherein any available position may be independently substituted with one or more R-groups or one or more activating moieties as described above.
where M, X, and k, are as defined above.

In some embodiments, at least one activating moiety is tethered to the diamine-derived portion of the salen ligand, as shown in formula X:

$$\text{X}$$

where M, X, k, R', and (Z), are as defined above.

In certain embodiments, salen ligands of formula X are selected from an optionally substituted moiety consisting of:

$$\text{Xa}$$

$$\text{Xb}$$

$$\text{Xc}$$

$$\text{Xd}$$

$$\text{Xe}$$

$$\text{Xf}$$

where M, X, k, R', and (Z), are as defined above.
where M, X, k, R', and \( (Z)_n \) are as defined above.

In certain embodiments, the diamine bridge of catalysts of formula Xa an optionally substituted moiety selected from the group consisting of:

where M and \( (Z)_n \) is as defined above.

In certain embodiments, metallosalenate complexes of the present invention include, but are not limited to those in Table 1 below:
TABLE 1 (continued)

\[
\begin{align*}
\text{N} & \quad \text{N} \\
\text{N} & \quad \text{N} \\
\text{N} & \quad \text{N} \\
\text{N} & \quad \text{N} \\
\text{N} & \quad \text{N} \\
\text{N} & \quad \text{N} \\
\text{N} & \quad \text{N} \\
\end{align*}
\]
<table>
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<th>N</th>
<th>N</th>
<th>21 NY</th>
<th>N</th>
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</thead>
</table>

TABLE 1 (continued)

![Chemical Structures](image-url)
In certain embodiments, for complexes of Table 1, M is Co—X, where X is as defined above. In certain embodiments, for complexes of Table 1, M is Co—OC(O)CF₃. In certain embodiments, for complexes of Table 1, M is Co—OAc. In certain embodiments, for complexes of Table 1, M is Co—OC(O)CF₃. In certain embodiments, for complexes of Table 1, M is Co—N₃. In certain embodiments, for complexes of Table 1, M is Co—Cl. In certain embodiments, for complexes of Table 1, M is Co-nitrophenoxo. In certain embodiments, for complexes of Table 1, M is Co-dinitrophenoxo.

In some embodiments, for complexes of Table 1, M is Cr—X, where X is as defined above.

In certain embodiments, a tetradeutate ligand is a porphyrin ligand. In some embodiments, a metal complex is a cobalt porphyrin complex. In certain embodiments, a metal complex is a chromium porphyrin complex. In some embodiments, a metal complex is an aluminum porphyrin complex.

Examples of porphyrin containing metal complexes of the present invention include, but are not limited to:
wherein each of M, X, k, R', and ——— (Z)m is as defined above.

In certain embodiments, a multidentate ligand is an optionally substituted tetrabenzo porphyrin. Suitable examples include, but are not limited to:
wherein M, R', and \( (Z)_n \) are as previously defined.

In certain embodiments of porphyrin and phthalocyanine-based complexes described herein, M is aluminum. In certain embodiments of porphyrin and phthalocyanine-based complexes described herein, M is cobalt. In certain embodiments of porphyrin and phthalocyanine-based complexes described herein, M is manganese.

In certain embodiments, porphyrin complexes of the present invention include, but are not limited to those in Table 2 below:
In certain embodiments, for complexes of Table 2, M is Co—X, where X is as defined above. In certain embodiments, for complexes of Table 2, M is Co—O(C=O)CF₃. In certain embodiments, for complexes of Table 2, M is Co—OAc. In certain embodiments, for complexes of Table 2, M is Co—Nₓ. In certain embodiments, for complexes of Table 2, M is Co—Cl. In certain embodiments, for complexes of Table 2, M is Co—nitrophenoxy. In certain embodiments, for complexes of Table 2, M is Co-dinitrophenoxy.

In certain embodiments, for complexes of Table 2, M is Al—X, where X is as defined above. In certain embodiments, for complexes of Table 2, M is Cr—X, where X is as defined above.

In certain embodiments, porphyrin complexes of the present invention are synthesized as shown in the following schemes:
In some embodiments, the present disclosure provides methods of polymerization comprising contacting an epoxide with carbon dioxide in the presence of a provided metal complex to form a polycarbonate. In some embodiments, the present invention provides a method of polymerization, the method comprising:

a) providing an epoxide of formula:

![Diagram](image)

wherein:

- $R_d$ is hydrogen or an optionally substituted radical selected from the group consisting of $C_{1-30}$ aliphatic; $C_{1-12}$ heteroaromatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and each of $R_o$, $R_e$, and $R_d$ is independently hydrogen or an optionally substituted radical selected from the group consisting of $C_{1-12}$ aliphatic; $C_{1-12}$ heteroaromatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

b) contacting the epoxide and carbon dioxide in the presence of a metal complex as described herein to provide a polymer having a formula selected from the group consisting of:

![Diagram](image)

In some embodiments, a provided polymer has a formula:

![Diagram](image)

In some embodiments, carbon dioxide is optional and a provided polymer has a formula:

![Diagram](image)

In certain embodiments, $R_o$, $R_e$, and $R_d$ are each hydrogen. In certain embodiments, $R_d$ is optionally substituted $C_{1-12}$ aliphatic. In some embodiments, $R_e$ is optionally substituted $C_{1-12}$ heteroaromatic. In some embodiments, the epoxide is ethylene oxide, propylene oxide, or cyclohexene oxide.

In certain embodiments, one of $R_o$, $R_e$, and $R_d$ is hydrogen. In certain embodiments, two of $R_o$, $R_e$, and $R_d$ are hydrogen. In certain embodiments, three of $R_o$, $R_e$, and $R_d$ are hydrogen.

In certain embodiments, $R_o$ is hydrogen. In certain embodiments, $R_e$ is hydrogen. In certain embodiments, $R_d$ is hydrogen. In certain embodiments, $R_d$ is hydrogen.

In certain embodiments, $R_o$, $R_d$, $R_e$, and $R_d$ are each independently an optionally substituted $C_{1-30}$ aliphatic group.

In certain embodiments, $R_o$, $R_d$, $R_e$, and $R_d$ are each independently $C_{1-12}$ heteroaromatic group. In certain embodiments, $R_o$, $R_d$, $R_e$, and $R_d$ are each independently an optionally substituted $C_{1-30}$ aliphatic group.

In certain embodiments, $R_o$, $R_d$, $R_e$, and $R_d$ are each independently an optionally substituted $C_{1-12}$ heteroaromatic group. In certain embodiments, $R_o$, $R_d$, $R_e$, and $R_d$ are each independently an optionally substituted $C_{1-30}$ aliphatic group.

In certain embodiments, $R_o$, $R_d$, $R_e$, and $R_d$ are each independently an optionally substituted $C_{1-12}$ aliphatic group. In certain embodiments, $R_o$, $R_d$, $R_e$, and $R_d$ are each independently an optionally substituted $C_{1-30}$ aliphatic group.

In certain embodiments, $R_d$ is an optionally substituted $C_{1-30}$ aliphatic group. In certain embodiments, $R_d$ is an optionally substituted $C_{1-12}$ aliphatic group. In certain embodiments, $R_d$ is an optionally substituted $C_{1-30}$ aliphatic group. In certain embodiments, $R_d$ is an optionally substituted $C_{1-12}$ aliphatic group. In certain embodiments, $R_d$ is an optionally substituted $C_{1-30}$ aliphatic group.
ally substituted 3-12-membered carboxyclic rings. In some embodiments, an \( R' \) and an \( R'' \) attached to the same carbon are taken together to form a polycyclic carboxyclic comprising two or more optionally substituted 3-8-membered carboxyclic rings. In some embodiments, an \( R' \) and an \( R'' \) attached to the same carbon are taken together to form a polycyclic carboxyclic comprising two or more optionally substituted 5-7-membered carboxyclic rings.

In some embodiments, an \( R' \) and an \( R'' \) attached to the same carbon are taken together to form a bicyclic carboxyclic comprising two optionally substituted 3-12-membered carboxyclic rings. In some embodiments, an \( R' \) and an \( R'' \) attached to the same carbon are taken together to form a bicyclic carboxyclic comprising two optionally substituted 3-8-membered carboxyclic rings. In some embodiments, an \( R' \) and an \( R'' \) attached to the same carbon are taken together to form a bicyclic carboxyclic comprising two optionally substituted 5-7-membered carboxyclic rings.

In certain embodiments, an \( R' \) and an \( R'' \) attached to the same carbon are taken together to form an optionally substituted 3-12-membered carboxyclic ring. In certain embodiments, an \( R' \) and an \( R'' \) attached to the same carbon are taken together to form an optionally substituted 3-8-membered carboxyclic ring. In certain embodiments, an \( R' \) and an \( R'' \) attached to the same carbon are taken together to form an optionally substituted 5-7-membered carboxyclic ring.

In some embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form one or more optionally substituted 3-12-membered carboxyclic rings. In some embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form a polycyclic carboxyclic comprising two or more optionally substituted 3-8-membered carboxyclic rings. In some embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form a polycyclic carboxyclic comprising two or more optionally substituted 5-7-membered carboxyclic rings.

In some embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form a bicyclic carboxyclic comprising two optionally substituted 3-12-membered carboxyclic rings. In some embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form a bicyclic carboxyclic comprising two optionally substituted 3-8-membered carboxyclic rings. In some embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form a bicyclic carboxyclic comprising two optionally substituted 5-7-membered carboxyclic rings.

In certain embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form an optionally substituted 3-12-membered carboxyclic ring. In certain embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form an optionally substituted 3-8-membered carboxyclic ring. In certain embodiments, an \( R' \) and an \( R'' \) attached to adjacent carbon are taken together to form an optionally substituted 5-7-membered carboxyclic ring.

In certain embodiments, the polymer comprises a copolymer of two different repeating units where \( R' \), \( R'' \), and \( R''' \) of the two different repeating units are not all the same. In some embodiments, a polymer comprises a copolymer of three or more different repeating units wherein \( R' \), \( R'' \), and \( R''' \) of each of the different repeating units are not all the same as \( R' \), \( R'' \), and \( R''' \) of any of the other different repeating units. In some embodiments, a polymer is a random copolymer. In some embodiments, a polymer is a tapered copolymer.

In some embodiments, a polymer contains a metal complex as described herein. In some embodiments, a polymer comprises residue of a metal complex as described herein. In some embodiments, a polymer comprises a salt of an organic cation and \( X \), wherein \( X \) is a nucleophile or counterion. In some embodiments, \( X \) is 2,4-dinitrophenolate anion.

In some embodiments, \( R'' \) is optionally substituted \( C_{1-12} \) aliphatic. In some embodiments, \( R'' \) is optionally substituted \( C_{1-12} \) heterocyclic. In some embodiments, \( R'' \) is optionally substituted phenyl. In some embodiments, \( R'' \) is optionally substituted 8- to 10-membered aryl. In some embodiments, \( R'' \) is optionally substituted 5- to 10-membered heterocyclic. In some embodiments, \( R'' \) is optionally substituted 3- to 7-membered heterocyclic.

In certain embodiments, \( R'' \) is selected from methyl, ethyl, propyl, butyl, vinyl, allyl, phenyl, trifluoromethyl.
In certain embodiments, $R'$ is

![Chemical structure](image)

In certain embodiments, $R''$ is

![Chemical structure](image)

In certain embodiments, $R'''$ is

![Chemical structure](image)

In some embodiments, $R''$ is hydrogen. In some embodiments, $R''$ is optionally substituted C$_{1-12}$ aliphatic. In some embodiments, $R''$ is optionally substituted C$_{1-12}$ heteroaliphatic. In some embodiments, $R''$ is optionally substituted phenyl. In some embodiments, $R''$ is optionally substituted 8- to 10-membered aryl. In some embodiments, $R''$ is optionally substituted 5- to 10-membered heteroaryl. In some embodiments, $R''$ is optionally substituted 3- to 7-membered heterocyclic.

In some embodiments, $R''$ is hydrogen. In some embodiments, $R''$ is optionally substituted C$_{1-12}$ aliphatic. In some embodiments, $R''$ is optionally substituted C$_{1-12}$ heteroaliphatic. In some embodiments, $R''$ is optionally substituted phenyl. In some embodiments, $R''$ is optionally substituted 8- to 10-membered aryl. In some embodiments, $R''$ is optionally substituted 5- to 10-membered heteroaryl. In some embodiments, $R''$ is optionally substituted 3- to 7-membered heterocyclic.

In some embodiments, $R''$ and $R'''$ are taken together with intervening atoms to form one or more rings selected from the group consisting of: optionally substituted C$_3$-C$_{14}$ carbocycle, optionally substituted 3- to 14-membered heterocycle, optionally substituted phenyl, optionally substituted C$_8$-C$_{10}$ aryl, and optionally substituted 5- to 10-membered heteroaryl.

In some embodiments, $R''$ and $R'''$ are taken together with intervening atoms to form one or more rings selected from the group consisting of: optionally substituted C$_3$-C$_{14}$ carbocycle, optionally substituted 3- to 14-membered heterocycle, optionally substituted phenyl, optionally substituted C$_8$-C$_{10}$ aryl, and optionally substituted 5- to 10-membered heteroaryl.

**EXAMPLES**

**Example 1**

A general route to a symmetric cobalt (III) salen ligand of the present invention is shown in Schemes E1 and E2, below:
As shown in Scheme E1, disubstituted phenol E1-a is formylated to provide salicylaldehyde derivative E1-b. Two equivalents of this aldehyde are then reacted with a diamine (in this case 1,2-diamino cyclohexane) to afford Schiff base E1-c. This compound is then reacted with cobalt (II) acetate to give the Co(II)-salen complex (not shown) which is oxidized by air in the presence of trifluoroacetic acid to afford the active cobalt (III) catalyst. Similar chemistries can be applied to synthesis of the catalysts described hereinabove. One skilled in the art of organic synthesis can adapt this chemistry as needed to provide the specific catalysts described herein.

**Example 2**

A typical route to an asymmetric cobalt (III) salen ligand is shown in Scheme E2:
As shown in Scheme E2, disubstituted salicylaldehyde derivative E1-b is treated with one equivalent of a monohydrochloride salt of 1,2-cyclohexanediamine the resulting Schiff base E2-a is then neutralized and a second different salicylaldehyde derivative is added. This compound is then reacted with cobalt (II) acetate to give the Co(II)-salen complex which is oxidized by air in the presence of trifluorooracetic acid to afford the active cobalt (III) catalyst. Similar chemistries can be applied to synthesis of the catalysts described herein above. One skilled in the art of organic synthesis can adapt this chemistry as needed to provide the specific catalysts described herein.

**Example 3**

Example 3 describes the synthesis of a catalyst

![Scheme E3](image)

where M is Co(III), is salicyl, is a P-linked phosphorimine moiety and m is 1, wherein there are one or two (Z)ₙ groups present (Scheme E4 and E3, respectively).

![Diagram](image)
As shown in Scheme E3, triol E3-a is protected as a ketal to afford monohydric alcohol E3-b, this compound is then alkylated with bromide E3-c to afford benzylic ether E3-d. Deprotection and oxidation of the other benzylic alcohol affords salicylaldehyde E3-e which is condensed with cyclohexanediamine as described above to give ligand E3-f. The phosphorimine nitrogen is then quaternized and the metal complex formed as before to provide catalyst E3-h. In an alternative route not shown here the metal is first inserted and then quarternization is performed.

Scheme E3b
As shown in Scheme E3b, salicylaldehyde E3-e (described above) is condensed with cyclohexanediamine monohydrochloride to afford the mono-Schiff base hydrochloride E4a. This salt is then neutralized, condensed with di-t-butyl salicylaldehyde, and methylated to give E4-b. The resulting ligand is metallated and oxidized as described above for Scheme E3 to give catalyst E4-c.

Example 4

Example 4 describes the synthesis of catalysts where M is Co(III),
Scheme E4 shows the synthesis of compounds CS-6 and CS-7. For each compound trans-1,2-Diaminocyclohexane (2.0 mol) is slowly added to an anhydrous ethanol solution of benzyl chloride CS-4 (1.0 mol). The reaction is stirred and heated to reflux for 3 h, then cooled to rt and diluted with water. This mixture is cooled overnight in the freezer and solids are collected by filtration to afford dichloride CS-5. The dichloride CS-5 (1.0 mol) is reacted with N,N-Dimethylamino pyridine (2.0 mol) or N-methyl imidazole in acetonitrile. The reactions are heated at 80°C for 18 h and then the solvent is removed in vacuo to provide the respective ammonium salts. These salts are metallated and oxidized as described previously to provide catalysts CS-6 and CS-7.

**Example 5**

Example 5 describes the synthesis of catalysts where M is Co(III), 

\[ Z \text{ is a } 1\{-[N\text{-methylimidazolium}] \text{ (CS-8), or dimethylamino (CS-9) and } m \text{ is 1, wherein there are two } (Z)_{\text{m}} \text{ groups present (Scheme E5 and E6, respectively).} \]

Scheme E5 shows the synthesis of compounds CS-8 and CS-9 using conditions similar to those described above. Synthesis of CS-8: The known compound 1-(2-methylaminoethyl)-3-methylimidazole (2.0 mol) is combined with CS-5 (1.0 mol) in acetonitrile. The reaction is heated to 80°C for 18 h and then the solvent is removed in vacuo, metallation with Co(OAc)₃ and oxidation in TFA are then performed as described above to afford catalyst CS-8. Synthesis of CS-9: N,N,N Trimethyl-1,2-ethanediame (4.0 mol) is combined with CS-5 (1.0 mol) in acetonitrile. The reaction is heated to 80°C for 18 h, cooled, and the solvent is removed in vacuo. The crude product is diluted with ether, filtered to remove amine salts, and concentrated in vacuo. The residue is dissolved in degassed methanol and combined with Co(OAc)₃ (1.0 mol). After stirring for 3 h the residue is filtered and washed with methanol. Trifluoroacetic acid (1.0 mol) is added slowly to a stirring solution of the solid residue in dichloromethane. After stirring open to air for 3 h, the solids are filtered and dried in vacuo to produce CS-9.

**Example 6**

Example 6 and Scheme E6 describe the synthesis of catalysts where M is Co(III), 

\[ Z \text{ is dibutylamino and } m \text{ is 1, wherein there are two } (Z)_{\text{m}} \text{ groups present.} \]
Synthesis of CS-10:
Ligand CS-5 (1.0 mol), 3-(dibutylamino)-1-propanol (2.0 mol), a 50% NaOH solution (10 mol), tetrabutylammonium bisulphate (4 mol %), and dichloromethane are combined and heated at 65°C overnight. The reaction mixture is concentrated in vacuo to remove the bulk of the solvent and the aqueous layer is extracted with ethyl acetate. The organic layer is separated, dried with magnesium sulfate, filtered, and concentrated in vacuo. After purification using silica gel the product is dissolved in degassed methanol and combined with Co(OAc)₃ (1.0 mol). After stirring for 3 h, the residue is filtered and washed with methanol. Trifluoroacetic acid (1.0 mol) is added slowly to a dichloromethane solution of the solid residue. After stirring open to air for 3 h, the solids are filtered and dried in vacuo to produce CS-10.

Example 7

Example 7 and Scheme E7 describe the synthesis of catalysts where M is Co(III),

is salicy,  ——— includes two

is salicy, ——— is

groups taken together to form a ring including the Z group, Z is 3-[N-methylpyridinium] and m is 1, wherein there is one ——— (Z)ₘ group present.

Z is 1-[4-t-butylpyridinium], and m is 2, wherein there are two ——— (Z)ₘ groups present.
Synthesis of AC-2.
Intermediate AC-1 (0.37 g, 0.35 mmol), 4-t-butylpyridine (0.21 mL, 1.41 mmol), and AcCN (4 mL) were combined in a sealed vial and heated to 80°C with stirring for 18 h. The solvent was removed in vacuo, leaving a yellow residue (0.61 g, 110% yield, AcCN present). 1H NMR (400 MHz, CDCl3, δ): 9.53 (t, 8H), 8.21 (s, 2H), 7.94 (t, 8H), 7.08 (s, 2H), 6.83 (s, 2H), 4.81 (m, 8H), 3.29 (m, 2H), 2.78 (m, 2H), 2.15 (s, 6H), 1.5-2.0 (m, 24H), 1.36 (s, 36H); IR (ATR, film cast from AcCN): ν_C=O=1637 cm⁻¹. A solution of the residue (0.30 g, 0.19 mmol) in dry EtOH (5 mL) was added to AgBF₄ (0.19 g, 0.85 mmol) in a schlenk tube and stirred overnight shielded from the light. The solution was filtered through Celite and the solvent was removed in vacuo, giving a solid residue. This residue was flushed over a small plug of silica gel with 5:1 CH₂Cl₂:EtOH as eluant. The solvent was removed to give a solid residue (0.18 g, 67% yield). 1H NMR (400 MHz, CDCl₃, δ): 8.75 (t, 8H), 7.98 (d, 2H), 7.92 (t, 8H), 7.1-7.3 (m, 4H), 4.52 (m, 8H), 3.6 (m, 21H), 2.7 (m, 2H), 2.19 (s, 6H), 1.5-2.0 (m, 24H), 1.38 (s, 36H); IR (ATR, film cast from CH₂Cl₂): ν_C=O=1641 cm⁻¹, ν_BF₄=1050 cm⁻¹. A solution of the residue (0.18 g, 0.12 mmol) in dry EtOH (4 mL) was added to Co(OAc)₃ (0.022 g, 0.12 mmol) in a schlenk tube under N₂. The solution was stirred for 3 h at room temperature, and the solvent was removed in vacuo. The residue was triturated with ether, dried in vacuo, and redissolved in CH₂Cl₂. A solution of CF₃CO₂H (9 μL, 0.12 mmol) in CH₂Cl₂ (80 μL) was added and the solution stirred for 3 h open to air. Solid Na₂O₂CCF₃ (0.067 g, 0.49 mmol) was added, and the solution was stirred under N₂ for 2 days. The solution was filtered through Celite and the solvent was removed in vacuo to leave a brown residue (0.071 g, 37% yield). 1H NMR (400 MHz, DMSO-d₆, δ): 8.86 (t, 8H), 8.08 (t, 8H), 8.07 (s, 2H), 7.30 (m, 4H), 4.44 (m, 8H), 3.54 (m, 2H), 2.9 (m, 2H), 2.47 (s, 6H), 1.5-2.0 (m, 24H), 1.29 (s, 36H); IR (ATR): ν_C=O=1682 cm⁻¹, ν_C=O=1641 cm⁻¹.

Additional ligands AC-6 through AC-11 were synthesized using the conditions described for compound AC-2 and are summarized in Scheme E8b and Table E8.
Example 9 and Scheme E9 describe the synthesis of catalysts where M is Co(III),

is salicyl, is

Z is N,N-bis-(3-dimethylaminopropyl)amino (AC-4), tetramethyl guanidino (AC-5), N-linked morpholino (AC-6), or N-linked piperidino (AC-14), and m is 2, wherein there are two \((Z)_{2m}\) groups present.
Synthesis of AC-4.
Intermediate AC-3 (0.45 g, 0.63 mmol), 3,3'-iminobis(N,N'-dimethylpropylamine) (0.28 mL, 1.26 mmol), K$_2$CO$_3$ (0.55 g, 2.52 mmol) and AcCN (5 mL) were combined in a sealed vial and heated to 80°C with stirring for 18 h. The solution was filtered and the solvent was removed in vacuo, triturated with ether, and dried in vacuo to leave a yellow residue (0.48 g, 91% yield). 1H NMR (400 MHz, CDCl$_3$, δ): 8.21 (m, 2H), 6.8-7.2 (m, 4H), 3.75 (m, 2H), 3.0-3.4 (m, 2H), 2.0-2.8 (m, 2H), 2.18 (s, 6H), 1.4-2.0 (m, 24H); IR (ATR): ν$_{C=O}$=1600 cm$^{-1}$. A solution of the residue (0.21 g, 0.25 mmol) in dry EtOH (10 mL) was added to Co(OAc)$_2$ (0.045 g, 0.25 mmol) in a schlenk tube under N$_2$, CH$_2$Cl$_2$ (5 mL) was added to completely dissolve the solution. The solution was stirred for 18 h at room temperature, and the solvent was removed in vacuo. The residue was triturated with ether, dried in vacuo, and redisolved in CH$_2$Cl$_2$ (10 mL). CF$_3$CO$_2$H (20 μL, 0.25 mmol) was added and the solution stirred for 3.5 h open to air. The solvent was removed in vacuo, triturated with ether, and dried in vacuo to leave a brown residue (0.28 g, 108% yield, residual CH$_2$Cl$_2$). 1H NMR (400 MHz, DMSO-d$_6$, δ): 7.92 (m, 2H), 7.1-7.4 (m, 4H), 3.58 (m, 2H), 3.0-3.4 (m, 20H), 2.0-2.8 (m, 28H), 2.3 (s, 6H), 1.4-2.0 (m, 24H); IR (ATR): ν$_{C=O}$=1688 cm$^{-1}$, ν$_{C=N}$=1616 cm$^{-1}$.

Synthesis of AC-5.
Intermediate AC-3 (0.20 g, 0.29 mmol), 1,1,3,3-tetramethylguanidine (0.21 mL, 1.71 mmol), K$_2$CO$_3$ (0.39 g, 2.85 mmol) and AcCN (2 mL) were combined in a sealed vial and reacted as in AC-4, except that the residue was also washed with hexanes. 1H NMR (400 MHz, CDCl$_3$, δ): 8.23 (d, 2H), 6.95 (s, 2H), 6.80 (s, 2H), 3.35 (m, 2H), 3.1 (m, 4H), 2.7-2.8 (m, 2H), 2.48 (m, 4H), 2.20 (s, 6H), 1.4-2.0 (m, 16H); IR (ATR): ν$_{C=O}$=1594 cm$^{-1}$. The residue was reacted as in AC-4. IR (ATR): ν$_{C=O}$=1690 cm$^{-1}$, ν$_{C=N}$=1610 cm$^{-1}$.

Synthesis of AC-6.
Intermediate AC-3 (0.32 g, 0.44 mmol), morpholine (0.16 mL, 1.77 mmol), K$_2$CO$_3$ (0.61 g, 4.4 mmol) and AcCN (4 mL) were combined in a sealed vial and reacted as in AC-4, except that the residue was also washed with a NaOAc buffer (pH=4) solution to remove residual morpholine. 1H NMR (400 MHz, CDCl$_3$, δ): 8.22 (s, 2H), 6.92 (s, 2H), 6.79 (s, 2H), 3.69 (m, 8H), 3.28 (m, 2H), 2.2-2.5 (m, 16H), 2.19 (s, 6H), 1.4-2.0 (m, 16H). The residue was reacted as in AC-4. 1H NMR (400 MHz, DMSO-d$_6$, δ): 7.91 (s, 2H), 7.23 (s, 2H), 7.14 (s, 2H), 3.6 (m, 2H), 3.52 (m, 8H), 2.99 (m, 2H), 2.94 (s, 2H), 2.47 (m, 4H), 2.2-2.5 (m, 12H), 1.4-2.0 (m, 16H); IR (ATR): ν$_{C=O}$=1671 cm$^{-1}$, ν$_{C=N}$=1630 cm$^{-1}$.
Additional ligands AC-13 and AC-14 were synthesized using the conditions described for compounds AC-4 through AC-6 and are summarized in Scheme E9b and Table E9:

![Scheme E9b](image)

**TABLE E9**

<table>
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<tr>
<th>Compound</th>
<th>Q (Scheme E9b)</th>
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<th>IR* (cm⁻¹)</th>
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<td>8.25 (s, 2H), 7.02 (d, 2H), 6.77 (d, 2H), 3.7 (m, 8H), 3.3 (m, 2H), 2.2-2.6 (m, 12H), 1.3-2.0 (m, 20H), 1.37 (s, 18H)</td>
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<tr>
<td>AC-14</td>
<td><img src="image" alt="Structure" /></td>
<td>ν(C=O) = 1629</td>
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</table>

*I400 MHz, CDCl₃
*All compounds exhibited the loss of a peak at 1213 cm⁻¹ attributed to the CH₃ group in AC-1.

Example 10

Confirmation of inventive concepts, processes, methods, and compositions described herein has been provided, among other ways, through publication by others after the priority date of the present case. For example, Examples 10-27 describe working Examples presented in Chinese Patent Application No. 200810229276.1, published as CN 101412809A. Additional experimental and characterization data are described by Lu and co-workers, J. Am. Chem. Soc., 2009, 131, 11509-11518, and supporting information available at www.pubs.acs.org, the entirety each of which is hereby incorporated by reference.

In certain embodiments, provided catalysts and/or methods for the preparation of polycarbonate are characterized by one or more of the following: retaining high catalytic activity at low catalyst concentration; reaction conditions that are relatively mild; high catalytic activity with high selectivity for polymer product; alternate structure in the polycarbonate product higher than 97% with relatively narrow distribution of molecular weight; retaining high catalytic activity for copolymerization of carbon dioxide and epoxides at higher reaction temperatures (e.g., above 50°C, above 75°C, or above 100°C); and catalysts that can be used to catalyze the polymerization of carbon dioxide with two or more alkylene oxides for the synthesis of polycarbonate polymer.

The following materials were added sequentially into a stainless steel high pressure reactor of effective volume of 200 mL at ambient temperature: 0.1 mmole of cobalt complex I-a (R₁ is cyclohexyl diamine, X is NO₃⁻¹ anion; R₂=H; R₃, R₄ and R₅ are tertiary butyl; group containing organic base group is at position 3 of the benzene ring in the ligand, n is 2) and one mole of propylene oxide. The reactor was then filled with carbon dioxide and the pressure is maintained constant at 2.0 MPa. The temperature was controlled at 25°C. The content was stirred with a magnetic stirring bar for 6 hours and the remaining carbon dioxide was slowly released. The remaining alkylene oxide was collected in -20°C. Cold trap and a certain amount of mixture of methanol/chloroform was added to dissolve the high polymer. Then a large amount of diethyl ether was added to precipitate the polycarbonate. The precipitate was filtered and washed several times with diethyl ether and dried in vacuum to constant weight to afford 27 grams of polycarbonate as a white solid. The average molecular weight of the polymer was determined by gel permeation chromatography to be 101,000 g/mol with a molecular weight distribution of 1.24. A Varian INOVA-400 MHz Nuclear Magnetic Resonance spectrometer was used to determine its ¹H-NMR and the result showed that the alternate structure is over 99%.
Example 11

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the molar ratio of catalyst to propylene oxide was changed from 1:100000 to 1:500000 (0.02 mmole of catalyst and 1 mole of propylene oxide were used). The reaction was carried out at 25°C for 24 hours to afford 21 grams of poly(propylene carbonate) with a molecular weight of 223,000 g/mol and a molecular weight distribution of 1.29. The polymer formed contained more than 99% carbonate linkages.

Example 12

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the molar ratio of catalyst to propylene oxide was changed from 1:100000 to 1:200000 (0.008 mmole of catalyst and 1.6 mole of propylene oxide were used). The reaction was carried out at 50°C for 10 hours to afford 19 grams of poly(propylene carbonate) with a molecular weight of 318,000 g/mol and a molecular weight distribution of 1.37. The polymer formed contained more than 99% carbonate linkages.

Example 13

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the molar ratio of catalyst to propylene oxide was changed from 1:10000 to 1:2000 (0.5 mmole of catalyst and 1 mole of propylene oxide were used). The reaction was carried out at 25°C for 3 hours to afford 48 grams of poly(propylene carbonate) with a molecular weight of 52,800 g/mol and a molecular weight distribution of 1.30. The polymer formed contained more than 99% carbonate linkages.

Example 14

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the reaction temperature was changed from 25°C to 100°C and the reaction was carried out for 0.5 hours to afford 34 grams of poly(propylene carbonate) with a molecular weight of 112,400 g/mol and a molecular weight distribution of 1.38. The polymer formed contained more than 99% carbonate linkages.

Example 15

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the reaction temperature was changed from 25°C to 10°C and the reaction was carried out for 10 hours to afford 18 grams of poly(propylene carbonate) with a molecular weight of 914,000 g/mol and a molecular weight distribution of 1.38. The polymer formed contained more than 99% carbonate linkages.

Example 16

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the propylene oxide was replaced with 1,2-butylene oxide. The reaction was carried out at 25°C for 6 hours to afford 31 grams of poly(butylene carbonate) with a molecular weight of 127,000 g/mol and a molecular weight distribution of 1.21. The polymer formed contained more than 99% carbonate linkages.

Example 17

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the propylene oxide was replaced with 1,2-octylene oxide. The reaction was carried out at 25°C for 10 hours to afford 34 grams of poly(octylene carbonate) with a molecular weight of 109,000 g/mol and a molecular weight distribution of 1.38. The polymer formed contained more than 99% carbonate linkages.

Example 18

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the propylene oxide was replaced with a mixture of propylene oxide and cyclohexene oxide (the molar ratio of the catalyst to propylene oxide and cyclohexene oxide was 1:5000:5000). The reaction was carried out at 50°C for 6 hours to afford 59 grams of poly(propylene-co-cyclohexene carbonate) with a molecular weight of 187,000 g/mol and a molecular weight distribution of 1.29. The polymer formed contained more than 99% carbonate linkages.

Example 19

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the axial anion in the cobalt complex 1-a was changed from nitrate radical to acetate moiety. The reaction was carried out at 25°C for 6 hours to afford 34 grams of poly(propylene carbonate) with a molecular weight of 95,000 g/mol and a molecular weight distribution of 1.28. The polymer formed contained more than 99% carbonate linkages.

Example 20

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the diamine skeleton in the cobalt
complex 1-a was changed from cyclohexane diamine to ethylene diamine. The reaction was carried out at 25°C for 6 hours to afford 29 grams of poly(propylene carbonate) with a molecular weight of 112,000 g/mol and a molecular weight distribution of 1.20. The polymer formed contained more than 99% carbonate linkages.

Example 21

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the diamine skeleton in the cobalt complex 1-a was changed from cyclohexane diamine to o-phenylene diamine. The reaction was carried out at 25°C for 6 hours to afford 25 grams of poly(propylene carbonate) with a molecular weight of 92,000 g/mol and a molecular weight distribution of 1.15. The polymer formed contained more than 99% carbonate linkages.

Example 22

The following materials were added sequentially into a stainless steel high pressure reactor of volume of 200 mL at ambient temperature: 0.1 mmole of cobalt complex 1-b (R₁ is 1,2-propylene diamine, X is di-nitrophenyl anion; R₂ = H; R₃ is tertiary butyl; there are organic base groups at position 5 of the two benzene rings in the ligand; n is 0) and 1 mole of propylene oxide. The reactor was then filled with carbon dioxide and the pressure was maintained constant at 2.0 MPa. The reaction was carried out at 25°C for 6 hours to afford 23 grams of polycarbonate as a white solid. The average molecular weight of the polymer was determined by gel permeation chromatography to be 81,000 g/mol with a molecular weight distribution of 1.34. The polymer formed contained more than 99% carbonate linkages.

Example 23

The following materials were added sequentially into a stainless steel high pressure reactor of volume of 200 mL at ambient temperature: 0.1 mmole of cobalt complex 1-c (R₁ is ethylene diamine, X is dinitrophenyl anion; R₂ = H; R₃ is tertiary butyl; there are organic base groups at position 3 and position 5 of one of the benzene rings in the ligand; n is 0) and 1 mole of propylene oxide. The reactor was then filled with carbon dioxide and the pressure was maintained constant at 2.0 MPa. The reaction was carried out at 25°C for 6 hours to afford 23 grams of polycarbonate as a white solid. The average molecular weight of the polymer was determined by gel permeation chromatography to be 81,000 g/mol with a molecular weight distribution of 1.34. The polymer formed contained more than 99% carbonate linkages.

Example 24

The following materials are added sequentially into a stainless steel high pressure reactor of volume of 200 mL at ambient temperature: 0.1 mmole of cobalt complex 1-b (R₁ is ethylene diamine, X is dinitrophenyl anion; R₂ = H; R₃ and R₄ are tertiary butyl; there are organic base groups at position 5 of the two benzene rings in the ligand; n is 0) and 1 mole of propylene oxide. The reactor was then filled with carbon dioxide and the pressure was maintained constant at 2.0 MPa. The reaction was carried out at 25°C for 6 hours to afford 26 grams of polycarbonate as a white solid. The average molecular weight of the polymer was determined by gel permeation chromatography to be 83,000 g/mol with a molecular weight distribution of 1.19. The polymer formed contained more than 99% carbonate linkages.

Example 25

The following materials are added sequentially onto a stainless steel high pressure reactor of volume of 200 mL at ambient temperature: 0.1 mmole of cobalt complex 1-a (R₁ is 2,3-butylene diamine, X is nitrate anion; R₂ = H; R₃, n and R₄ are methoxyl group; R₅ is tertiary butyl; there is an organic base group at position 3 of one of the benzene rings in the ligand; n is 2) and 1 mole of propylene oxide. The reactor was then filled with carbon dioxide and the pressure is maintained constant at 2.0 MPa. The reaction was carried out at 25°C for 6 hours to afford 22 grams of polycarbonate as a white solid. The average molecular weight of the polymer was determined by gel permeation chromatography to be 73,000 g/mol with a molecular weight distribution of 1.14. The polymer formed contained more than 99% carbonate linkages.
Example 26

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the pressure was changed from 2.0 MPa to 0.1 MPa. The reaction was carried out at 25°C for 10 hours to afford 25 grams of poly(propylene carbonate) with a molecular weight of 100,400 g/mol and a molecular weight distribution of 1.17. The polymer formed contained more than 99% carbonate linkages.

Example 27

The same equipment and reaction conditions were employed as in Example 10 with the same catalyst and the same conditions except that the pressure was changed from 2.0 MPa to 6.0 MPa. The reaction was carried out at 25°C for 6 hours to afford 29 grams of poly(propylene carbonate) with a molecular weight of 125,000 g/mol and a molecular weight distribution of 1.25. The polymer formed contained more than 99% carbonate linkages.

While we have described a number of embodiments of this invention, it is apparent that our basic examples may be altered to provide other embodiments that utilize the compounds and methods of this invention. Therefore, it will be appreciated that the scope of this invention is to be defined by the appended claims rather than by the specific embodiments that have been presented by way of example.

What is claimed is:

1. A metal complex for the synthesis of aliphatic poly carbonate polymers comprising a cobalt salen complex, wherein the salen ligand has an activating moiety tethered to the position ortho to a metal-bound oxygen substituent of one or both of the salicylaldehyde-derived phenyl rings of the salen ligand,

   wherein each activating moiety has a formula \( \text{----(Z)---} \), and comprises a linker \( \text{-----} \) coupled to at least one activating functional group \( Z \), with \( m \) denoting the number of activating functional groups present on a single linker moiety; \( m \) is an integer from 1 to 4;

---

### Table E10

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### Results

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<tr>
<td>18</td>
<td>24</td>
<td>25.5</td>
<td>2547</td>
<td>424.4</td>
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<tr>
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<td>25</td>
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<td>2255</td>
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the linker comprises an optionally substituted C_{2-30} aliphatic group wherein one or more methylene units are optionally and independently replaced by —NR', —N(R')C(O)—, —C(O)N(R')—, —O—, —C(O)—, —OC(O)—, —C(O)O—, —S—, —SO—, —SO_2—, —C(=S)—, —C(C=N)—, or —N—N—; where each occurrence of R'y is independently —H, or an optionally substituted radical selected from the group consisting of C_{1-6} aliphatic, 3- to 7-membered heterocyclic, phenyl, and 8- to 10-membered aryl; and Z comprises a neutral nitrogen-containing functional group, a cationic functional group, or an arsonium functional group.

2. The metal complex of claim 1, wherein the cobalt salen complex comprises a substructure independently selected from the group consisting of:

3. The metal complex of claim 1, wherein Z comprises a neutral nitrogen-containing functional group.

4. The metal complex of claim 1, wherein the neutral nitrogen-containing functional group comprises a guanidine or amidine.

5. The metal complex of claim 3, wherein the neutral nitrogen-containing functional group is selected from the group consisting of:
or a combination of two or more of these, wherein:

R₁ and R₂ is independently at each occurrence hydrogen or an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein an R₃ group can be taken with an R¹ or R² group to form one or more optionally substituted rings; and

R¹ is independently each occurrence, hydrogen, a hydroxyl protecting group, or an optionally substituted radical selected from the group consisting of C₁₋₂₀ alkyl; C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

6. The metal complex of claim 1, wherein the linker "-----" comprises an optionally substituted C₅₋₁₀ aliphatic group.

7. The metal complex of claim 1, wherein the linker "-----" comprises an optionally substituted C₄₋₆ aliphatic group.

8. The metal complex of claim 1, wherein the linker "-----" comprises an optionally substituted C₄ aliphatic group.

9. The metal complex of claim 1, wherein the linker "-----" comprises an optionally substituted C₁₃ aliphatic group.

10. The metal complex of claim 1, wherein the linker "-----" comprises an optionally substituted straight alkyl chain.

11. The metal complex of claim 1, wherein the linker "-----" is selected from the group consisting of:
where * represents the site of attachment to the salen ligand, and each # represents a site of attachment of an activating functional group; and

R¹ is independently at each occurrence, —H or an optionally substituted radical selected from the group consisting of C₆H₅, aliphatic, 3- to 7-membered heterocyclic, phenyl, and 8- to 10-membered aryl.

12. The metal complex of claim 1, wherein the cobalt has an oxidation state of 3⁺.

13. The metal complex of claim 1, wherein there are activating moieties tethered to the positions ortho and para to the phenolic oxygen of one or both of the salicylaldehyde-derived phenyl rings of a salen ligand.

14. The metal complex of claim 1, wherein m is 1.

15. The metal complex of claim 1, wherein m is greater than 1.

16. The metal complex of claim 1, wherein Z is a cationic functional group selected from the group consisting of:
or a combination of two or more of these, wherein:

R', R'', and R''' are independently at each occurrence hydrogen, or an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; wherein any two or more R', R'', and R''' groups can be taken together with intervening atoms to form one or more optionally substituted rings optionally containing one or more additional heteroatoms;

R'' is hydrogen or —OR'';

R''' is hydrogen, hydroxyl, or optionally substituted C₁₋₂₀ aliphatic;

R'' and R''' are independently at each occurrence, hydrogen or an optionally substituted radical selected from the group consisting of C₁₋₂₀ aliphatic; C₁₋₂₀ heteroaliphatic; phenyl; a 3- to 8-membered saturated or
partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered unsaturated monocyclic carbocycle; a 5- to 6-membered monocyclic saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocyclic ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; where each occurrence of R is independently —H, or an optionally substituted radical selected from the group consisting of C_1-C_6 aliphatic, 3- to 7-membered heterocyclic, phenyl, and 8- to 10-membered aryl, and where two or more adjacent R groups can be taken together to form an optionally substituted saturated, partially unsaturated, or aromatic 5- to 12-membered ring containing 0 to 4 heteroatoms; X is any anion, and

Ring A comprises an optionally substituted, 5- to 10-membered heteroaryl group.

17. The metal complex of claim 16, wherein the cationic moiety comprises a guanidinium group.

18. The metal complex of claim 16, wherein the cationic moiety comprises an amidinium group.

19. The metal complex of claim 16, wherein the linker "—R" comprises an optionally substituted C_5-C_30 aliphatic group.

20. The metal complex of claim 16, wherein the linker "—R" comprises an optionally substituted C_4-C_30 aliphatic group.

21. The metal complex of claim 16, wherein the linker "—R" comprises an optionally substituted C_6 aliphatic group.

22. The metal complex of claim 16, wherein the linker "—R" comprises an optionally substituted straight alkyl chain.

23. The metal complex of claim 16, wherein the linker "—R" comprises an optionally substituted straight alkyl chain.

24. The metal complex of claim 16, wherein the linker "—R" is selected from the group consisting of: 

[Diagram of molecular structures is shown]
where * represents the site of attachment to the salen ligand, and each # represents a site of attachment of an activating functional group; and

R is independently at each occurrence, —H or an optionally substituted radical selected from the group consisting of C1−8 aliphatic, 5- to 7-membered heterocyclic, phenyl, and 8- to 10-membered aryl.

25. The metal complex of claim 16, wherein the cobalt has an oxidation state of 3+.

26. The metal complex of claim 16, wherein there are activating moieties tethered to the positions ortho and para to the phenolic oxygen of one or both of the salicylaldehyde-derived phenyl rings of a salen ligand.

27. The metal complex of claim 16, wherein m is 1.

28. The metal complex of claim 16, wherein m is greater than 1.

29. The metal complex of claim 1, wherein Z is an arsionium group.

30. The metal complex of claim 29, wherein the arsionium group comprises:

where R, R', and R'' are, independently at each occurrence, hydrogen or an optionally substituted radical selected from the group consisting of C1−20 aliphatic; C1−20 heteroaliphatic; phenyl; a 3- to 8-membered saturated or partially unsaturated monocyclic carbocycle, a 7-14 carbon saturated, partially unsaturated or aromatic polycyclic carbocycle; a 5- to 6-membered monocyclic heteroaroyl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 3- to 8-membered saturated or partially unsaturated heterocyclic ring having 1-3 heteroatoms independently selected from nitrogen, oxygen, or sulfur; a 6- to 12-membered polycyclic saturated or partially unsaturated heterocycle having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur; and an 8- to 10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur;
wherein any two or more R^1, R^2 and R^3 groups can be
taken together with intervening atoms to form one or
more optionally substituted rings; and

X^- is any anion.

31. The metal complex of claim 30, wherein the arsonium
group is selected from the group consisting of:

32. A polymerization method comprising the step of con-
tacting an epoxide with carbon dioxide in the presence of an
effective amount of a metal complex of claim 3.

33. The polymerization method of claim 32, wherein the
neutral nitrogen-containing functional group is selected from
the group consisting of:

R^2 and R^3 is independently at each occurrence hydrogen, or
an optionally substituted radical selected from the group
consisting of C_1-20 aliphatic; C_1-20 heteroaliphatic; pheno-
lyl; a 3- to 8-membered saturated or partially unsatur-
ated monocyclic carbocycle, a 7-14 carbon saturated,
partially unsaturated or aromatic polycyclic carbocycle;
a 5- to 6-membered monocyclic heteroaryl ring having
1-4 heteroatoms independently selected from nitrogen,
oxogen, or sulfur; a 3- to 8-membered saturated or par-
tially unsaturated heterocyclic ring having 1-3 heteroa-
toms independently selected from nitrogen, oxygen, or sulfur;
a 6- to 12-membered polycyclic saturated or partially
unsaturated heterocyclic ring having 1-5 heteroatoms
independently selected from nitrogen, oxygen, or sulfur;
an 8- to 10-membered bicyclic heteroaryl ring hav-
ing 1-5 heteroatoms independently selected from nitrogen,
oxogen, or sulfur; wherein R^1 and R^2 groups can be
taken together with intervening atoms to form one or
more optionally substituted rings optionally containing
one or more additional heteroatoms;

R^3 is independently at each occurrence hydrogen, or
an optionally substituted radical selected from the group
consisting of C_1-20 aliphatic; C_1-20 heteroaliphatic; pheno-
lyl; a 3- to 8-membered saturated or partially unsatur-
ated monocyclic carbocycle, a 7-14 carbon saturated,
partially unsaturated or aromatic polycyclic carbocycle;
a 5- to 6-membered monocyclic heteroaryl ring having
1-4 heteroatoms independently selected from nitrogen,
oxogen, or sulfur; a 3- to 8-membered saturated or par-
tially unsaturated heterocyclic ring having 1-3 heteroa-
toms independently selected from nitrogen, oxygen, or sulfur;
a 6- to 12-membered polycyclic saturated or partially
unsaturated heterocyclic ring having 1-5 heteroatoms
independently selected from nitrogen, oxygen, or sulfur;
an 8- to 10-membered bicyclic heteroaryl ring hav-
ing 1-5 heteroatoms independently selected from nitrogen,
oxogen, or sulfur; wherein an R^3 group can be taken
with an R^1 or R^2 group to form one or more optionally
substituted rings; and

R^4 is independently each occurrence, hydrogen, a hydroxyl
protecting group, or an optionally substituted radical
selected from the group consisting of C_1-20 acyl; C_1-20
aliphatic; C_1-20 heteroaliphatic; phenyl; a 3- to 8-mem-
bered saturated or partially unsaturated monocyclic car-
bocycle, a 7-14 carbon saturated, partially unsaturated
or aromatic polycyclic carbocycle; a 5- to 6-membered
monocyclic heteroaryl ring having 1-4 heteroatoms
independently selected from nitrogen, oxygen, or sulfur;
a 3- to 8-membered saturated or partially unsaturated
heterocyclic ring having 1-3 heteroatoms independently
selected from nitrogen, oxygen, or sulfur; a 6- to
12-membered polycyclic saturated or partially unsatur-
ated heterocyclic ring having 1-5 heteroatoms indepen-
dently selected from nitrogen, oxygen, or sulfur; and an 8- to
10-membered bicyclic heteroaryl ring having 1-5 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

34. The polymerization method of claim 32, wherein the epoxide is selected from the group consisting of: ethylene oxide, propylene oxide, butylene oxide, cyclohexene oxide, 1,2-oxide, 3-vinyl cyclohexene oxide, epichlorohydrin and mixtures of any two or more of these.

35. The polymerization method of claim 32, wherein the epoxide is ethylene oxide, propylene oxide, or cyclohexene oxide.

36. The polymerization method of claim 32, wherein the epoxide is ethylene oxide.

37. The polymerization method of claim 32, wherein the epoxide is propylene oxide.

38. A polymerization method comprising the step of contacting an epoxide with carbon dioxide in the presence of an effective amount of a metal complex of claim 16.

39. The polymerization method of claim 38, wherein the epoxide is selected from the group consisting of: ethylene oxide, propylene oxide, butylene oxide, cyclohexene oxide, 1,2-oxide, 3-vinyl cyclohexene oxide, epichlorohydrin and mixtures of any two or more of these.

40. The polymerization method of claim 38, wherein the epoxide comprises ethylene oxide, propylene oxide, or cyclohexene oxide.

41. The polymerization method of claim 38, wherein the epoxide comprises ethylene oxide.

42. The polymerization method of claim 38, wherein the epoxide comprises propylene oxide.

* * * * *