A process is disclosed for producing CF3CH2F from CF3CHClF by catalytic chlorination of CF3CHClF to CF3CCl2F at elevated temperature, and reaction of CF3CCl2F with H2 in the presence of a carbon-supported precious metal catalyst at a temperature of from about 100°C to 250°C Suitable catalysts for the chlorination include carbon catalysts and catalysts wherein halides of certain metals (La, Zn, Cu, Cr, Ru, Rh, and/or Pt) are supported on carbon.

5606085

THERMALLY ACTIVATED OLEFIN METATHESIS CATALYST PRECURSOR

Bell Andrew; Coffy Tim West Grove, PA, UNITED STATES assigned to Metton America Inc

Molybdenum and tungsten compounds that are useful as catalyst precursors in the metathesis of olefins have the general formula:(*See Patent for Tabular Presentation*) PS wherein M is tungsten or molybdenum; Y is oxygen or NR1; R1, R2, and R3 are the same or different and are selected from alkyl, cycloalkyl, cycloalkenyl, polycycloalkyl, polycycloalkenyl, haloalkyl, haloaralkyl, substituted or unsubstituted aralkyl and aryl groups, and silicon-containing analogs thereof; L is a Lewis base; X is halogen; s is 0 or 1; x+y+z=4, and y> or =1, provided that when x is 2 or more, two OR2 groups can be replaced by a chelating ligand (OR2)2. These compounds can be used for the metathesis of olefins in neat metathesizable olefin, as well as in solution, and require only the input of energy to be converted to active catalysts.

5606103

ORGANOTIN CATALYZED TRANSESTERIFICATION

Trapasso Louis E; Meisel Philip L; Meisel Lee B; Chwang Willy K West Long Branch, NJ, UNITED STATES assigned to CPS Chemical Company Inc

Methyl or ethyl esters of mono- and polycarboxylic acids are transesterified with alcohols and polyols by reaction in the presence of a catalytically effective amount of organotin catalyst, so that an alcohol or polyol carboxylic acid ester is formed, which is then washed with aqueous alkali having a pH bigger than about 13.2 so as to remove essentially all of the organotin catalyst, thereby permitting the recovery of the alcohol or polyol carboxylic acid ester essentially free of the organotin catalyst.

5608096

METHOD OF FORMING SILOXANE POLYMERS USING A HETEROPOLY CATALYST HAVING A KEGGIN STRUCTURE

Katsoulis Dimitris; Keryk John R Midland, MI, UNITED STATES assigned to Dow Corning Corporation

The invention relates to a method of forming siloxane polymers using a heteropoly catalyst having a Keggin structure. The method comprises contacting a fluid comprising at least one siloxane polymer precursor selected from the group consisting of cyclic siloxanes (I) having the formula (RR'SiO)a and linear siloxanes (II) having the formula (*See Patent for Chemical Structure*) where R is a substituted or unsubstituted monovalent hydrocarbon having from 1 to 6 carbon atoms, R' is hydrogen or a substituted or

unsubstituted monovalent hydrocarbon having from 1 to 6 carbon atoms, a is at least 3 and b is at least 1; with an effective amount of a heteropoly catalyst having a Keggin structure, selected from the group consisting of heteropoly acids (III) having the formula HnXM12O40, salts thereof and mixed acid-salts thereof, where X is B+3. Si+4, Ge +4, p+5, M is Mo+6 and n is 3, 4 or 5, as required for the valence of the heteropoly acid to equal 0, at a temperature ranging from about 23°C to the boiling point of the siloxane polymer precursor.

5608113

PROCESS FOR PREPARATION OF DIAMINES BY CATALYTIC AMINATION OF AMINOALCOHOLS

Becker Rainer; Menger Volkmar; Reif Wolfgang; Henne Andreas Bad Durkheim, GERMANY assigned to BASF Aktiengesellschaft

In a process for preparing deamines from aminoalcohols and nitrogen compounds selected from the group consisting of ammonia and primary and secondary amines at from 80° to 250°C and pressures of from 1 to 400 bar using hydrogen in the presence of a zirconium, copper, nickel catalyst, the catalytically active composition comprises from 20 to 85% by weight of oxygen-containing zirconium compounds, calculated as ZrO2, from 1 to 30% by weight of oxygen-containing compounds of copper, calculated as CuO, from 30 to 70% by weight of oxygen-containing compounds of nickel, calculated as NiO, from 0.1 to 5% by weight of oxygen-containing compounds of molybdenum, calculated as MoO3, and from 0 to 10% by weight of oxygen-containing compounds of aluminum and/or manganese, calculated as Al2O3 and MnO2 respectively.

5608123

PROCESS FOR CATALYTIC HYDRATION OF OLEFINS

Inoue Kaoru; Iwasaki Masao; Ueda Naohiro Yokohama, JAPAN assigned to Mitsui Toatsu Chemicals Inc

There is provided a process for reacting water and olefin such as ethylene or propylene under mild conditions in the presence of a polyorganosiloxane contaminating sulfonic acid groups to produce alcohol corresponding to the olefin with high yield and selectivity.

5610113

PROCESS FOR THE PREPARATION OF ALPHA-TOCOPHEROL DERIVATIVES, AND CATALYST

Matsui Makoto; Yamamoto Hisash Aichi, JAPAN assigned to Eisai Co Ltd

A process is provided for the preparation of an alpha-tocopherol derivatives which are useful as antisterile vitamins, hypolipidemic blood flow increasing agents, anti-cytosenility agents. antioxidants and the like. Catalysts are also provided. The alpha-tocopherol derivatives are represented by the following formula (VII): (*See Patent for Chemical Structure*) (VII) wherein n stands for 0 or an integer of from 1 to 5. The derivative can be industrially prepared by employing as catalyst a metal ion-exchanged montmorillonite, metal ion-exchanged bentonite or metal ion-exchanged saponite which is substituted with one metal ion selected from the group consisting of scandium, yttrium, lanthanide element, aluminium, iron, tin, copper, titanium, zinc, nickel, gallium or zirconium.