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Discovery of Asymmetric Autocatalysis and the Study on the Origin of Homochirality of Biomolecules

1. Introduction

It is well known that life on the Earth is composed of L-amino acids and D-sugars. Like left and right hands, L-amino acid and D-amino acids are mirror images and are not superimposable (Fig. 1). They are termed as enantiomers and are chiral. One of the most characteristic features of life is that many biomolecules are highly enantioenriched enantiomers. If proteins and enzymes contain D-amino acids in random manner, the proteins and enzymes will not operate in normal manner. If DNA and RNA contain L-sugars in a random manner, the normal helical structure will not be formed and DNA and RNA will not work. In usual chemical reactions, the probability of the formation of D and L enantiomers is one to one, and the products are racemic modifications which contain one to one ratio of D and L enantiomers. In sharp contrast, all of the living organisms on the Earth are composed of essentially only L-amino acids. In addition, DNA and RNA of all living organisms employ only D-sugars. The chiral homogeneity of biomolecules, i.e., homochirality, is one of the striking features of life and is considered to be closely related to the origin of life.

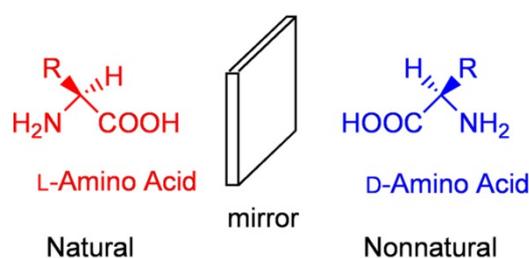


Fig. 1 Natural L-amino acid and its enantiomer.

Why, when and how did biomolecules become highly enantiomerically enriched? What is the origin of chirality? It has been considered that, on the early Earth or in space, by some reason, the initial racemic mixture of D and L enantiomers had become highly enantioenriched. Was it possible for such a reaction to really occur? Several theories have been proposed as the origins of chirality of organic compounds; right and left circularly polarized light (CPL), chiral inorganic crystals such as *d* and *l*-quartz, statistical fluctuation (spontaneous generation). However, the enantiomeric imbalances

induced by these mechanisms have been very low (less than 2% enantiomeric excess). Therefore, in order to reach the high enantioenrichment observed in nature, the process has been needed for significant amplification of extremely low enantiomeric excess to very high enantiomeric excess.

Soai discovered asymmetric autocatalysis in which chiral product acts as chiral catalyst for its own production with significant amplification of enantiomeric excess. Pyrimidyl alkanol acts as asymmetric autocatalyst in the enantioselective addition of diisopropylzinc to pyrimidine-5-carbaldehyde to afford more of itself with remarkable amplification of enantiomeric excess (>99.5% ee). Moreover, he found asymmetric autocatalysis initiated by chiral quartz, circularly polarized light, chiral crystal formed from achiral organic compounds as the origins of chirality. In addition, he reported the first experimental example of a spontaneous absolute asymmetric synthesis by using asymmetric autocatalysis with amplification of chirality. The reaction found by Soai has been often mentioned as the Soai reaction and Soai Asymmetric Autocatalysis. [1]

2. Discovery of Asymmetric Autocatalysis

Asymmetric autocatalysis, i.e., asymmetric automultiplication and catalytic replication of chiral compound, is a reaction in which the structures of chiral catalyst and the product are the same and in which chiral product acts as a chiral catalyst for its own production (Fig. 2). In usual asymmetric catalysis, because the structures of chiral catalyst and the product are different, it is necessary to separate the product from the catalyst. In addition, because the amount of catalyst doesn't increase during the reaction, the amount of the recovered catalyst decreases and the catalytic activity intrinsically decrease.

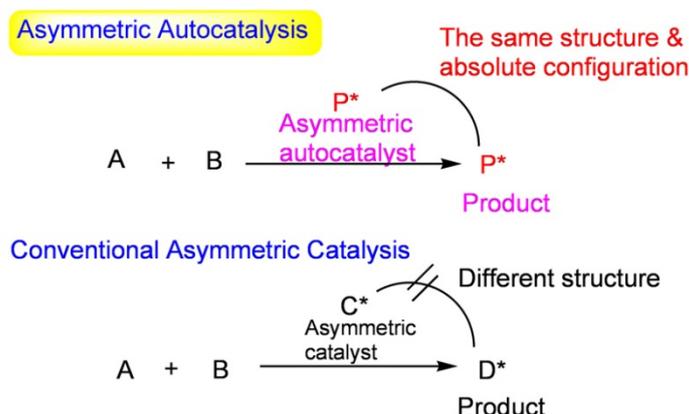


Fig. 2 Asymmetric autocatalysis and conventional asymmetric catalysis.

Unlike usual asymmetric catalysis, asymmetric autocatalysis has the following advantages: Because the structures of catalyst and the product are the same, it's not

necessary to separate the product from the catalyst. The efficiency of the reaction is high, because the amount of the catalyst increases during the reaction. The decrease in the amount and the catalytic activity of chiral catalyst can be avoided. In asymmetric autocatalysis, it can be anticipated that the initial tiny imbalance of chirality can be amplified without the need of any other chiral compound.

Soai found unprecedented asymmetric autocatalysis of pyridyl alkanol [2] in 1990, pyrimidyl alkanol [3] in 1995 and quinolyl alkanol. Pyrimidyl alkanol acts as an asymmetric autocatalyst in the enantioselective addition of diisopropylzinc to pyrimidine-5-carbaldehyde to afford the same pyrimidyl alkanol of the same absolute configuration with the initial asymmetric autocatalyst (Fig. 3). [3] The reaction is an automultiplication of enantioenriched pyrimidyl alkanol. Near perfect asymmetric autocatalysis with greater than 99.5% enantiomeric excess has been achieved. [4] When the product, i.e., pyrimidyl alkanol, is used as an asymmetric autocatalyst for the next round, the enantiomeric excess is greater than 99.5%. Even after 10th round, the enantioselectivity of product & catalyst is greater than 99.5%, only one enantiomer has automultiplied in near quantitative yield. The result shows that, during the 10 consecutive rounds of asymmetric autocatalysis, the initial enantiomer of pyrimidyl alkanol has automultiplied by a factor of 60 million times.

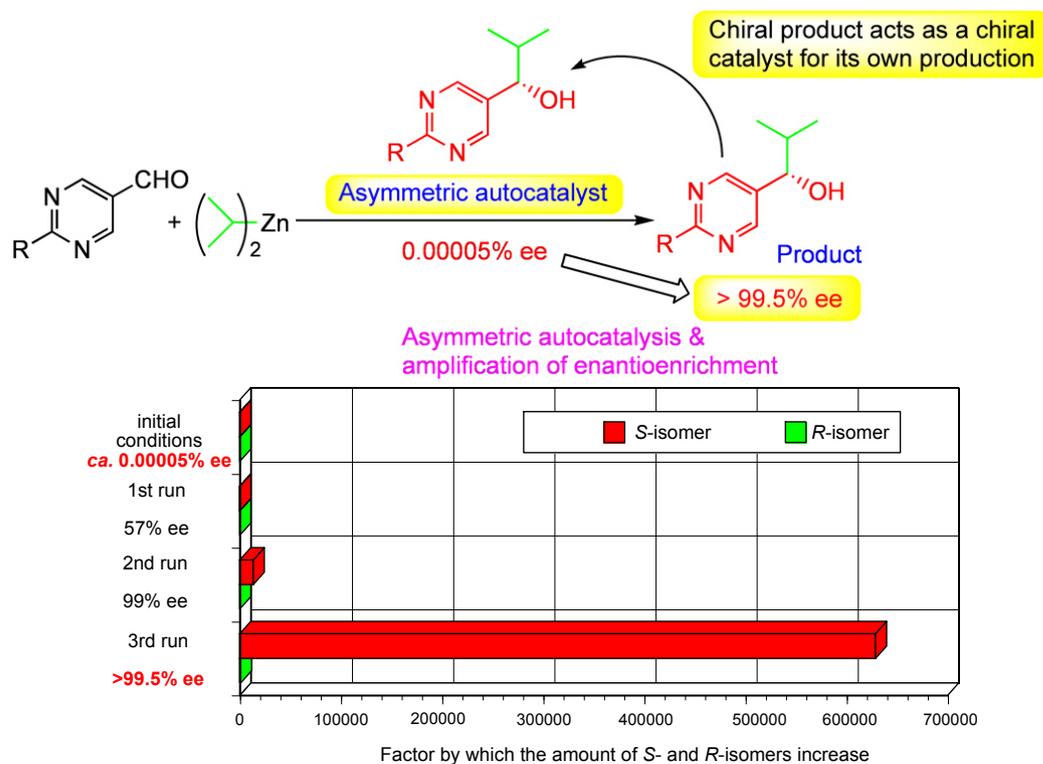


Fig. 3 Asymmetric autocatalysis with amplification of enantiomeric excess.

L-leucine.

Soai found that asymmetric autocatalysis can correlate the low enantioenrichment of the obtained leucine with the high enantioenrichment of organic compound. Thus, in the presence of L-leucine with low (2%) enantioenrichment, the reaction between pyrimidine-5-carbaldehyde and diisopropylzinc and the subsequent asymmetric autocatalysis afforded highly enantioenriched pyrimidyl alkanol with the correlated absolute configuration with amino acid. [6] The initially formed pyrimidyl alkanol has tiny enantiomeric excess owing to the presence of enantioenriched leucine, and the subsequent asymmetric autocatalysis with amplification of chirality affords the highly enantioenriched pyrimidyl alkanol. It is reported that strong CPL occurs in the star formation region in space.

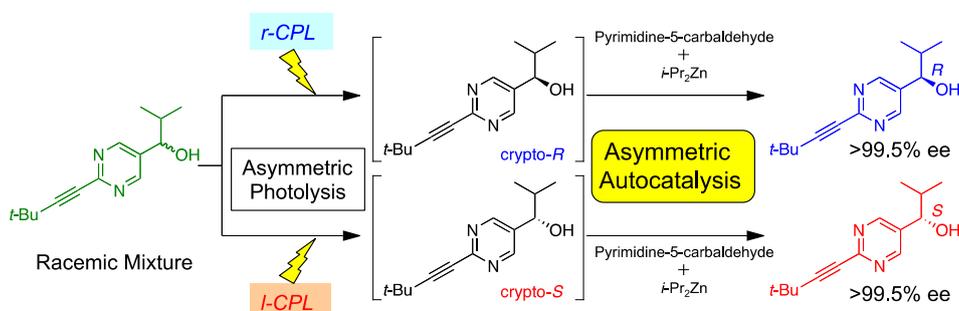


Fig. 5 Asymmetric autocatalysis triggered by circularly polarized light.

Direct irradiation of left-CPL (wavelength: 313 nm) to racemic (mixture of R and S enantiomers with the ratio of 50:50) pyrimidyl alkanol, and the subsequent asymmetric autocatalysis affords (*S*)-pyrimidyl alkanol with greater than 99.5% enantiomeric excess (fig. 5). [7] On the contrary, the irradiation with right-CPL (313 nm) and the asymmetric autocatalysis gave (*R*)-pyrimidyl alkanol with greater than 99.5% enantiomeric excess.

As described, asymmetric autocatalysis enables circularly polarized light (CPL) to play a role of the origin of chirality of organic compounds with very high enantiomeric excess.

3-2. Chiral Inorganic Crystals such as *d* and *l*-Quartz

Quartz occurs naturally and exhibits *d* and *l*-enantiomorphs. It has been considered that quartz is the origin of chirality of organic compounds. However, the enantiomeric excess induced by quartz has been extremely low and no experiment has been reported which correlates the chirality of quartz and that of organic compound with high enantiomeric excess. He found that, in the presence of the powder of *d*-quartz, the reaction between pyrimidine-5-carbaldehyde and diisopropylzinc and the subsequent

asymmetric autocatalysis affords (*S*)-pyrimidyl alkanol with very high enantiomeric excess, while in the presence of *l*-quartz (*R*)-alkanol with very high enantiomeric excess was formed (Fig. 6). [8] Thus, an organic compound with high enantiomeric excess is synthesized for the first time, in conjunction with asymmetric autocatalysis, using quartz as the origin of chirality. He also found that *d* and *l*-sodium chlorate serves as the chiral initiator of asymmetric autocatalysis. [9]

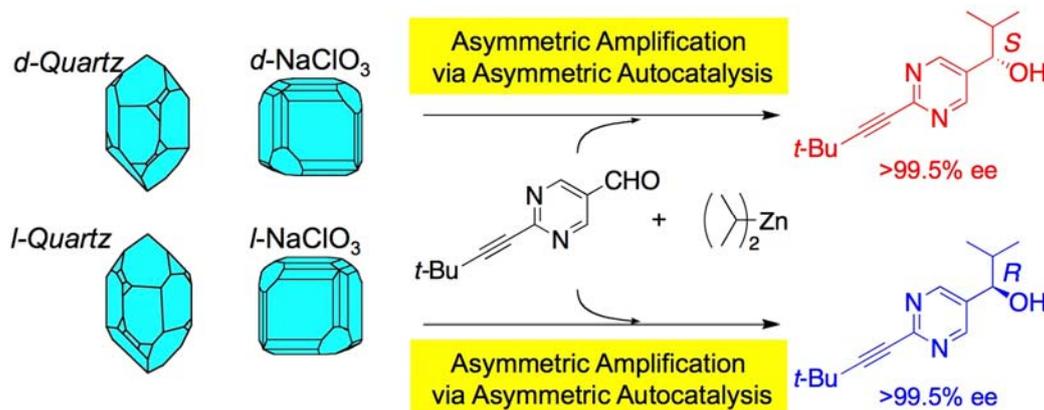


Fig. 6 Asymmetric autocatalysis triggered by chiral inorganic crystals.

As described, asymmetric autocatalysis with amplification of chirality afford, for the first time, the highly enantioenriched organic compound using quartz as the origin of chirality.

3-3. Spontaneous Absolute Asymmetric Synthesis

It has been well accepted that, without the intervention of any chiral factor, the probability of the formation of *R* and *S* product is fifty to fifty (50 : 50); racemate is formed. However, according to the theory of statistics, the numbers of *R* and *S* enantiomers are not exactly the same, i.e., there is almost always the fluctuation in numbers of enantiomers. Let's think about tossing coin in 100 times: the probability is only 0.08 (8 times) that you get exactly 50 heads and 50 tails in 100 times trials. In the rest of 0.92 probability (92 times), the numbers of head and tail are not the same. Moreover, when the total number is an odd number, you cannot get exactly the same number of head and tail. Like toss the coins, one can consider the initial fluctuation of chirality in organic reaction without the intervention of any chiral factor. Of course, when the number of molecules becomes large, the enantiomeric excess is below the detection level.

As described above, asymmetric autocatalysis of pyrimidyl alkanol has enormous power of amplification of enantiomeric excess from extremely low (*ca.* 0.00005% ee) to greater than 99.5% enantiomeric excess. [5]

Soai found that the reaction of pyrimidine-5-carbaldehyde with diisopropylzinc, without adding any chiral material, gives enantioenriched pyrimidyl alkanol with either *R* or *S* absolute configurations (Fig. 7). [10] The formation of *R* and *S* pyrimidyl alkanol exhibits approximate stochastic one to one distribution. It is conceivable that the initially formed stochastic fluctuation in chirality is amplified by asymmetric autocatalysis. The results of stochastic formation of *R* and *S* pyrimidyl alkanol fulfill the conditions necessary for spontaneous absolute asymmetric synthesis. This is the first example of a spontaneous absolute asymmetric synthesis. The result has significant implication in the origin and amplification of chirality in nature.

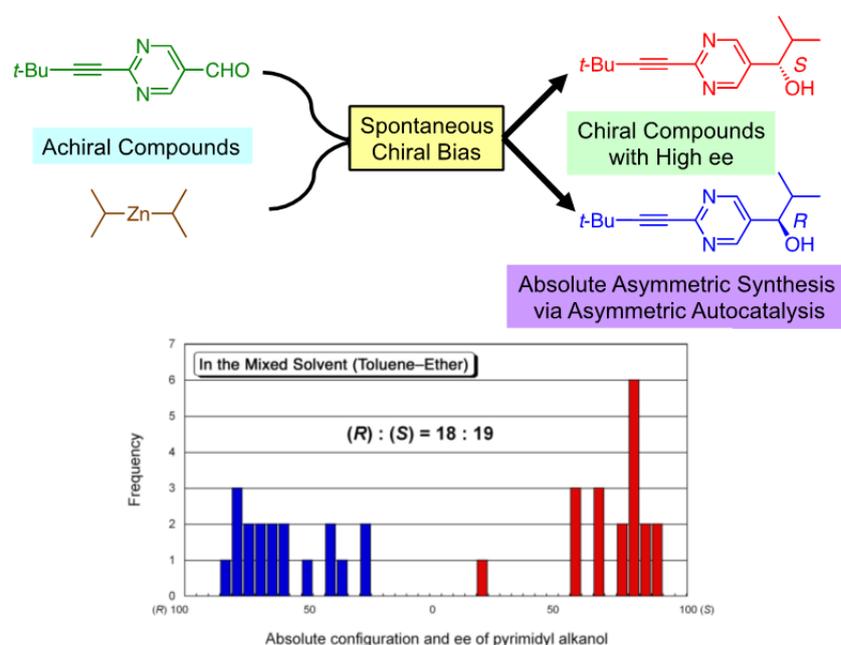


Fig. 7 Spontaneous absolute asymmetric synthesis.

3-4. Asymmetric Autocatalysis Initiated by Chiral Crystals formed from Achiral Organic Compounds

Some of the achiral organic compounds form chiral crystals. However, these chiral crystals have not been known to induce chirality in other compounds. He found that asymmetric autocatalysis is initiated by these chiral crystals formed from achiral organic compounds such as co-crystals and *N*-benzoylglycine (hippuric acid) to afford highly enantioenriched pyrimidyl alkanol with the corresponding absolute configurations with those of chiral organic crystals. [11a,b] Thus, by using asymmetric autocatalysis, the formation of chiral organic crystals from achiral organic compounds has been successfully correlated to high enantioenrichments of other organic compound.

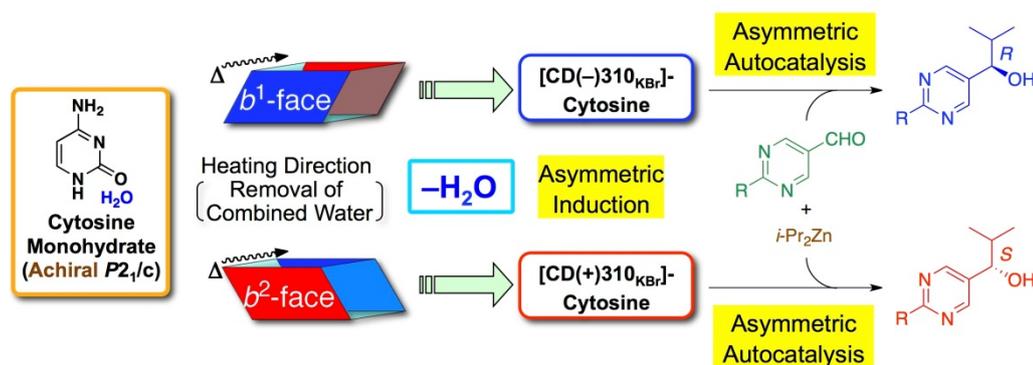


Fig. 8 Generation of crystal chirality of cytosine by dehydration from achiral crystal and asymmetric induction in the asymmetric autocatalysis.

In addition, he reported the formation of chiral cytosine crystals of desired chirality from the achiral cytosine hydrate crystal by dehydration from a specific face (fig. 8). [11c] Cytosine and adenine are achiral nucleobases, and have not been considered as the origin of chirality. He found chiral crystals of cytosine [11c] and adenine dinitrate [11d] act as chiral triggers of asymmetric autocatalysis to afford highly enantioenriched pyrimidyl alkanol with the corresponding absolute configurations to those of chiral crystals of cytosine and adenine dinitrate.

3-5. Asymmetric Induction by Chiral Compounds Arising from Isotope Substitution and by Cryptochiral Compounds.

Chiral isotopomers due to carbon ($^{13}\text{C}/^{12}\text{C}$) substitution have not been considered to act as chiral inducers in asymmetric reaction. Soai found that chiral compounds arising from carbon ^{13}C substitution act as chiral initiators and induce asymmetry in the addition of $i\text{-Pr}_2\text{Zn}$ to pyrimidine-5-carbaldehyde to afford highly enantioenriched pyrimidyl alkanol with the corresponding absolute configurations as those of the $^{13}\text{C}/^{12}\text{C}$ chiral compounds (Fig. 9).

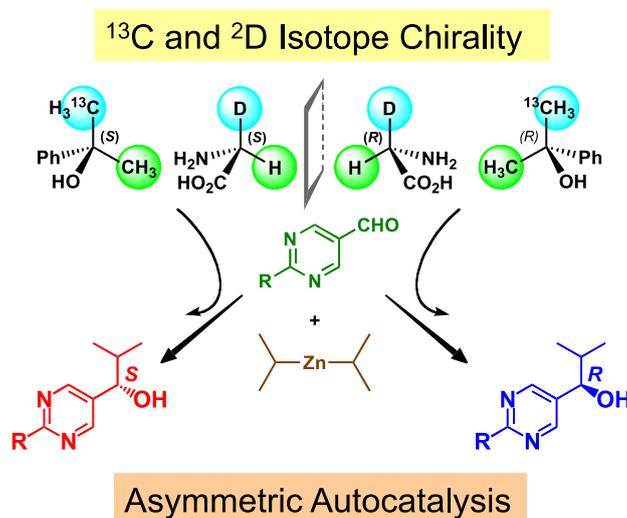


Fig. 9 Asymmetric autocatalysis triggered by chiral isotopomers.

[12a] He also found that chiral amino acids and primary alcohols as a result of deuterium substitution [12b] and chiral alcohol due to oxygen isotope substitution [12c] act as chiral initiators of asymmetric autocatalysis.

Butylethylhexylpropylmethane is a chiral saturated quaternary hydrocarbon. However, it is known that the compound does not exhibit any detectable optical rotation, thus the compound is cryptochiral. By using the cryptochiral compound, asymmetric autocatalysis was found to discriminate the chirality of the compound. [13] Cryptochiral isotactic polystyrene was also successfully discriminated.

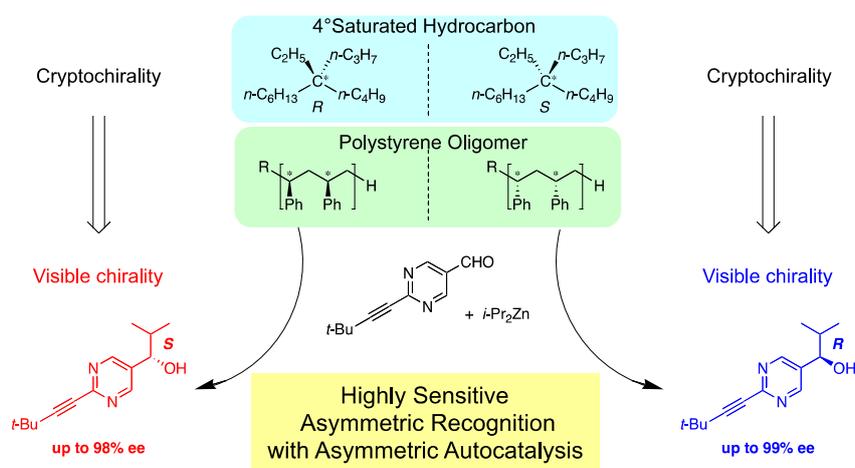


Fig. 10 Asymmetric autocatalysis induced by cryptochiral substances.

3-6. Asymmetric Induction on Enantiotopic Face of Achiral Crystal in Conjunction with Asymmetric Autocatalysis

Enantiotopic face of achiral crystal of certain pyrimidine-5-carbaldehyde was treated with vapor of diisopropylzinc to afford pyrimidyl alkanol with absolute configuration related to enantiotopic face. [14]

4. Summary

As described, Soai discovered asymmetric autocatalysis of pyridyl alkanol and pyrimidyl alkanol in the addition of diisopropylzinc to the corresponding pyridine-3-carbaldehyde and pyrimidine-5-carbaldehyde, respectively. Pyrimidyl alkanol was found to be practically perfect asymmetric autocatalyst, i.e., >99% yield and >99.5% ee. Moreover, asymmetric autocatalysis of pyrimidyl alkanol exhibit significant amplification of enantiomeric excess from very low (ca. 0.00005% ee) to near enantiopure (>99.5% ee) with significant amplification of quantity. He has proved that there is real chemical reactions in which very slight enantiomeric imbalance

becomes overwhelming by asymmetric autocatalysis. By using asymmetric autocatalysis, he found circularly polarized light, chiral inorganic crystal such as quartz, chiral crystal formed from achiral organic compound, chiral carbon isotopomers, cryptochiral compound such as saturated quaternary hydrocarbon trigger asymmetric autocatalysis to afford highly enantioenriched compound with the corresponding absolute configurations to those of chiral initiators. He also reported spontaneous absolute asymmetric synthesis using his asymmetric autocatalysis.

The asymmetric autocatalysis discovered by Soai has made a deep impact on scientific community in understanding the origin of chirality and homochirality. Moreover, the reaction is a very efficient method of catalytic asymmetric carbon-carbon bond forming reaction because of asymmetric autocatalysis. It has often been called as the Soai Reaction or Soai Asymmetric Autocatalysis.

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