Re-Engineering Nature's Catalysts
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fur might be associated with the radar anomalies. Neumann et al. did find this association, but also found the opposite—that the surfaces of the radar anomalies are not bright, but are typically quite dark relative to average Mercury reflectance with rare exceptions.

Paige et al. synthesized these observations to suggest that the dark deposits are due to organics more refractory than water ice, either directly deposited with the ice as part of the same process, or formed in situ by a low-temperature organic synthesis. Production of organics by irradiated cometary ices is well established in the laboratory and has been invoked to explain planetary and astrophysical observations of comets and the interstellar medium (see the figure). Paige et al. suggest that as a result of exposure to solar wind and ultraviolet radiation, complex organic radiolytic products are formed from ices and accumulate as a protective lag deposit that ultimately shields ice from further sublimation. They further note that dark deposits are observed to be more extensive than radar-bright material, and also occur in locations with somewhat higher model temperatures. This is evidence that ice deposits were more widespread in the past, and that the dark deposits in areas with higher temperature and lacking radar anomalies are relic deposits of this earlier era. A few restricted portions of the polar surface are substantially brighter than typical for Mercury, and Paige et al. show that these surfaces are cold enough to preserve surface frost against sublimation loss.

Prior to the MESSENGER results, polar ice at Mercury (and the Moon) was generally accepted, but the new data reveal a dynamic history of these deposits. The presence of the organic lag deposits strongly indicates that comets are the source of the polar volatiles, because other proposed sources are barren with respect to the critical elements needed for organic synthesis. The results also show that the charging of the cold traps can temporarily overcome thermal instability and can be used to derive a high lower limit on the amount of water vapor that can be at least transiently retained in a transient atmosphere of Mercury in a comet impact to account for the distribution of the dark deposits.

References

CHEMISTRY

Re-Engineering Nature’s Catalysts

Alison R. H. Narayan and David H. Sherman

Natural systems have inspired many scientific and technological advances (1). Materials design seeks to duplicate the fiber optical features of glass sponges (2); inorganic chemical complexes are modeled after enzyme active sites; and synthetic chemistry strategies parallel the biosynthetic pathways that produce complex natural product molecules (3). Scientists have also directly manipulated nature’s tools through enzyme and metabolic pathway engineering. On page 307 of this issue, Coelho et al. (4) report that a bacterial cytochrome P450—a protein that naturally catalyzes C–H bond oxidation—can be engineered to efficiently produce highly strained cyclopropanated products. This work demonstrates the biomimicry paradigm in reverse, where altering the function of one of nature’s most versatile biocatalysts is directed to a transformation originally conceived and implemented by synthetic chemists.

Throughout evolutionary time, enzymes have continuously morphed to perform specific functions on defined substrates in living cells. Advances in molecular biology have enabled new strategies to maximize the synthetic utility of particular enzymes and to expand their substrate scope and catalytic activity. Directed evolution is one of the most successful approaches toward achieving these goals (5). This method is a laboratory-based, accelerated version of natural evolution. Mutants with beneficial properties are advanced to subsequent rounds of mutagenesis in an iterative quest for the ideal enzyme. This approach has proven fruitful for altering the substrate scope for numerous target enzymes. For example, Arnold et al. evolved P450_{BM3} to selectively hydroxylate ethane, a much smaller molecule than the natural fatty acid substrate, to afford ethanol (6). Directed evolution has also been used to identify thermally stable enzyme mutants (7) or variants that function effectively in organic solvents (8).

These accomplishments have advanced the field of enzymatic catalysis. However, many continue to overlook the utility of enzymes compared to the endless modes of reactivity that can be probed and discovered by synthetic chemists. With a specific function ingrained in a protein’s native design, is it possible to completely re-engineer its function through iterative mutagenesis of its corresponding gene?

An engineered enzyme catalyzes a reaction that is fundamentally different from that catalyzed by the natural enzyme.

To access a totally new form of catalysis requires more than just library screening of a target enzyme. This feat demands a fundamental understanding of the enzyme’s structure and function combined with chemical intuition. In a pioneering study, Wilson and Whitesides used an oversized protein ligand to create a chiral environment around a metal complex. These artificial metalloenzymes, with biotin-tethered metal complexes bound to streptavidin, catalyzed asymmetric hydrogenation reactions (9). The power of this approach was recently expanded by Hyster et al., who used biotinylated rhodium(III) complexes with streptavidin mutants for asymmetric C–H activation. The study demonstrated the influence of key second sphere residues on the reactivity and selectivity of the enzyme (10).

Although progress has been made to optimize the native function of enzymes, the ability to leverage an individual active site to catalyze a new type of reaction remains a greater challenge. To achieve this goal, Coelho et al. selected diazoester carbene precursors as their cyclopropanation reagent. These compounds form metallacarbenoids with metals such as Rh, Ru, Cu, Co, and Fe (11). From the metallacarbenoid intermediate, cyclopropanation occurs through the transfer of the carbene to a double bond.

Coelho et al. screened 92 structurally and functionally diverse P450_{BM3} mutants from a larger library. They found five candidates,
Biomimicry in reverse. The natural P450 mono-oxygenase enzyme catalyzes C–H bond oxidation. Through judicious choice of enzyme and substrate followed by directed evolution, Coelho et al. have re-engineered this protein to perform a different catalytic reaction, namely cyclopropanation.

which they evolved further to a suite of di- stereo- and enantioselective cyclopropanation catalysts that can favor either a cis- or trans-cyclopropane product.

An existing library of P450<sub>BM3</sub> variants and information on the native functionality of these proteins facilitated this impressive example of engineering new enzyme reactivity. Pairing library screening with additional information on structure and function provides an appealing model for streamlined, directed evolution efforts. For example, Fasan et al. recently reported the development of P450 enzymes for the regio- and stereoselective oxidation of the human antimalarial drug artemisinin. The authors performed first-sphere active-site mutagenesis, followed by high-throughput screening with a panel of model compounds; they then used the reactivity data to predict substrate scope and to drive further protein evolution efforts (12).

These studies show that translating classical synthetic chemical transformations into versatile chemoenzymatic reactions is ripe for expansion. However, going from a promising idea on paper to an efficient enzymatic transformation remains a formidable challenge. Drawing on multiple sources of information—such as computational modeling, x-ray structural data, reactivity profiles, and high-density library design derived from iterative mutagenesis and screening—will be essential for building a wide variety of new functions into nature’s complex catalysts.

**References**

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**PERSPECTIVES**

**Polymer Rigidity Improves Microporous Membranes**

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Gas separation with membranes has been commercialized for more than 30 years, and includes processes such as the production of nitrogen (N<sub>2</sub>) from air and the removal of carbon dioxide (CO<sub>2</sub>) from natural gas. Commercial membranes have been largely derived from polymers with moderately rigid chains that pack closely to create small intermolecular spaces (or “free volume”) that impart moderate to high gas selectivity. However, their relatively low gas permeability slows down the separation processes. Microporous organic polymers (MOPs) (1–3) offer higher permeability, but the polymer chains must be made sufficiently rigid to maintain good selectivity. On page 303 of this issue, Carta et al. (4) describe a soluble, highly rigid MOP, from which a highly permeable membrane with good selectivity was fabricated. For example, oxygen (O<sub>2</sub>) and N<sub>2</sub> have only a 5% difference in kinetic diameters (which are related to the smallest effective dimensions of the gases), but the gas throughput of the smaller O<sub>2</sub> molecule is very much higher through their membrane.

Gases are separated by first pressurizing gas mixtures in contact with the membrane; each component sorbs (dissolves in it) and diffuses (passes through the free volume) through it at different rates with a character-