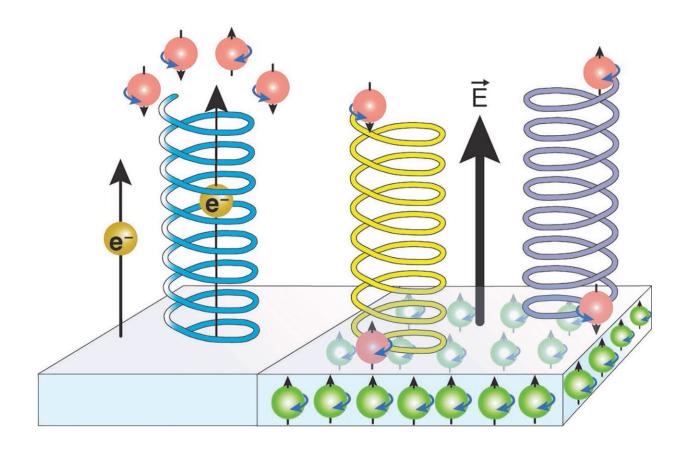


Magnetic effects at the origin of life? It's the spin that makes the difference

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Credit: Advanced Materials (2023). DOI: 10.1002/adma.202308666

Biomolecules such as amino acids and sugars occur in two mirror-image forms—in all living organisms, however, only one is ever found. Why this is the case is still unclear. Researchers at Empa and Forschungszentrum Jülich in Germany have now found evidence that the



interplay between electric and magnetic fields could be at the origin of this phenomenon.

The so-called homochirality of life—the fact that all biomolecules in living organisms only ever occur in one of two mirror-image forms—has puzzled a number of scientific luminaries, from the discoverer of molecular chirality, Louis Pasteur, to William Thomson (Lord Kelvin) and Nobel Prize winner Pierre Curie.

A conclusive explanation is still lacking, as both forms have, for instance, the same <u>chemical stability</u> and do not differ from each other in their physicochemical properties. The hypothesis, however, that the interplay between electric and magnetic fields could explain the preference for one or the other mirror-image form of a molecule—socalled enantiomers—emerged early on.

It was only a few years ago, though, that the first indirect evidence emerged that the various combinations of these force fields can indeed "distinguish" between the two mirror images of a molecule. This was achieved by studying the interaction of chiral molecules with <u>metallic</u> <u>surfaces</u> that exhibit a strong electric field over short distances.

The surfaces of magnetic metals such as iron, cobalt or nickel thus allow electric and magnetic fields to be combined in various ways—the direction of magnetization is simply reversed, from "North up—South down" to "South up—North down."

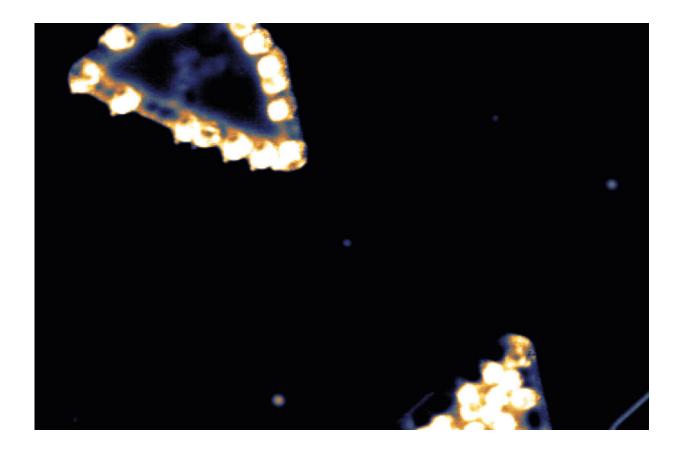
If the interplay between magnetism and electric fields actually triggers "enantioselective" effects, then the strength of the interaction between chiral molecules and magnetic surfaces should also differ, for example—depending on whether a right-handed or left-handed molecule "settles" on the surface.



Mirror images prefer opposing magnetic fields

And this is indeed the case, as a team of researchers led by Karl-Heinz Ernst from the Empa's Surface Science and Coating Technologies lab and colleagues at the Peter Grünberg Institute at the Forschungszentrum Jülich in Germany <u>reported</u> in the journal *Advanced Materials*.

The team coated a (non-magnetic) copper surface with small, ultra-thin "islands" of magnetic cobalt and determined the direction of the magnetic field in these using spin-polarized scanning tunneling microscopy; as mentioned before, this can run in two different directions perpendicular to the metal surface: North up or South up. They then deposited spiral-shaped chiral molecules—a 1:1 mixture of left- and right-handed heptahelicene molecules—onto these cobalt islands in ultrahigh vacuum.





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Then they "simply" counted the number of right- and left-handed helicene molecules on the differently magnetized cobalt islands, almost 800 molecules in total, again using scanning tunneling microscopy. And lo and behold: Depending on the direction of <u>magnetic field</u>, one or the other form of the helicene spirals had settled preferentially.

Moreover, the experiments showed that the selection—the preference for one or the other enantiomer—not only occurs during the binding on the cobalt islands, but already beforehand.

Before the molecules take up their final (preferred) position on one of the cobalt islands, they migrate long distances across the copper surface in a significantly weaker bound precursor state in "search" for an ideal position. They are only bound to the surface by so-called van der Waals forces. These are merely caused by fluctuations in the electronic shell of atoms and molecules and are therefore relatively weak. The fact that even these are influenced by magnetism, i.e. the direction of rotation (spin) of the electrons, was not known thus far.

Electrons with the 'wrong' spin are filtered out

Using scanning tunneling microscopy, the researchers were also able to solve another mystery, as they reported in the journal *Small* in November 2023. Electron transport—i.e. electric current—also depends on the combination of molecular handedness and magnetization of the surface.

Depending on the handedness of the bound molecule, electrons with one



direction of spin preferentially flow—or "tunnel"—through the molecule, meaning that electrons with the "wrong" spin are filtered out. This chirality-induced spin selectivity had already been observed in earlier studies, but it remained unclear whether an ensemble of molecules is necessary for this or whether individual molecules also exhibit this effect.

Ernst and his colleagues have now been able to show that individual helicene molecules also exhibit the CISS effect. "But the physics behind this is still not understood," admits Ernst.

The Empa researcher also believes that his findings eventually cannot fully answer the question of the chirality of life. In other words, the question that the Nobel Prize winner in chemistry and ETH chemist Vladimir Prelog described as "one of the first problems of molecular theology" in his Nobel Prize lecture in 1975.

But Ernst can imagine that in certain surface-catalyzed <u>chemical</u> <u>reactions</u>—such as those that could have taken place in the chemical "primordial soup" on the early Earth—a certain combination of electric and magnetic fields could have led to a steady accumulation of one form or another of the various biomolecules—and thus ultimately to the handedness of life.

More information: Mohammad Reza Safari et al, Enantioselective Adsorption on Magnetic Surfaces, *Advanced Materials* (2023). DOI: 10.1002/adma.202308666

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