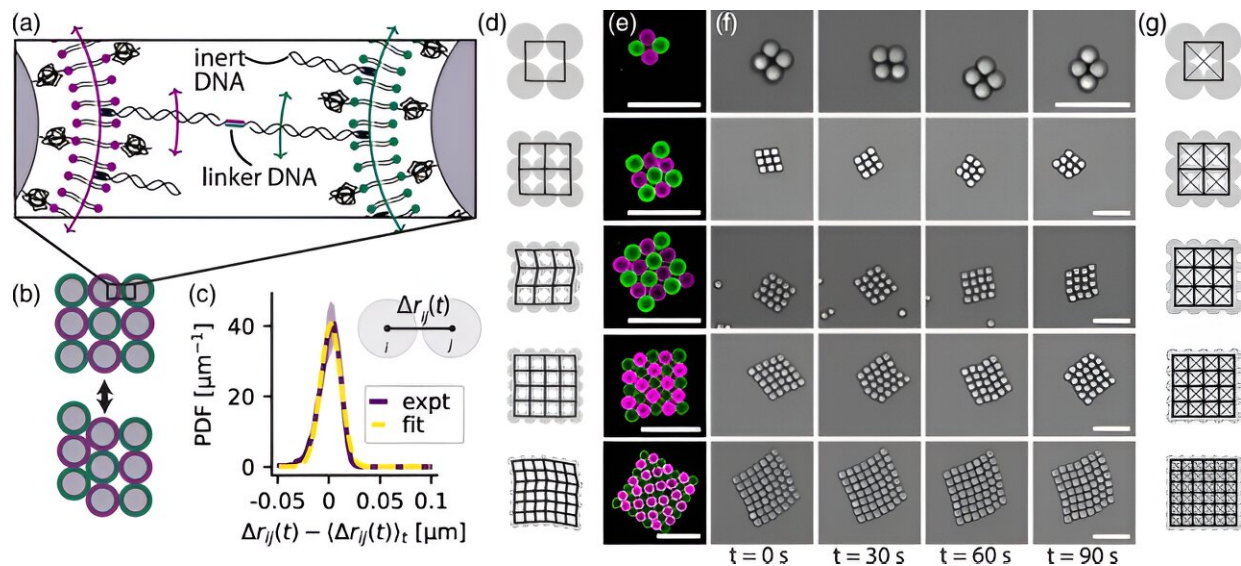


# A first in the lab: A tiny network of microparticles that is both strong and flexible

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Flexible colloidal square lattices. (a) A schematic overview of the binding mechanism. (b) A schematic example of a floppy mode deformation in a 3x3 square lattice. (c) The distribution of the distance between two particles. (d) Time-averaged particle positions as measured in experiments, where the bond network is indicated in black. (e) Confocal images of the  $n \times n$  square lattices. (f) Brightfield snapshots of the same lattices taken 30 s apart. (g) Spring networks used in the theoretical description. Credit: *Physical Review Letters* (2024). DOI: 10.1103/PhysRevLett.132.078202

Daniela Kraft's group has succeeded in creating a network of microparticles that is both strong and completely flexible. This may

sound simple, yet they are the first in the world to succeed in doing so. The achievement represents a real breakthrough in soft matter physics. [The study](#) is published in *Physical Review Letters*.

Ph.D. candidate Julio Melio studies microscopic, flexible networks and that's no easy job. In nature, such micro networks are found in gels, polymers or the cytoskeleton of the cells in your body. "These materials are pliable thanks to so-called soft modes, flexible states," Melio explains.

"We don't really know how temperature affects these states. It's too complicated to study this in [biological systems](#), so we made a [network](#) of microscopic spheres, colloids, in the lab. The simplest system is a square lattice. That can deform into a diamond-like shape, for example."

## **A clever technique for flexible connections**

The researcher buys silica colloids and gives them a coating of lipids. Then he creates a DNA link to connect the spheres. "We use two types of DNA strands that can attach to each other and place them on colloids. These can then bind to each other, but not to another [colloid](#) of the same species. The special thing about these DNA links is that the linked particles can move relative to each other. So the network is flexible."

Next begins the difficult job of getting the beads into the desired structure. That's quite a challenge, Melio explains. "You pick up one colloid with so-called optical tweezers, a laser, and put it in contact with a second one. That's how you build the lattice one by one." However, the system is extremely sensitive, so with the slightest change in circumstances you get qualitatively bad spheres that stick together. "And then the system loses its flexibility," says Melio.

The first time, it took the Ph.D. candidate nearly three-quarters of a year

to make a perfectly square grid of five by five colloids. "By now, I can fortunately do it a lot faster," he says. This makes Kraft's group the first in the world to build a large microstructure in such a controlled way without losing flexibility.

## Potential applications: Metamaterials and microrobots

The researchers have already gained new insights that help to better understand the soft modes in microgrids. The larger the lattice is, the more likely it is to be in the square state rather than the diamond one. Larger structures also shear better: they deform more easily under [shear force](#) than smaller variants.

This is interesting for developing new metamaterials, where the properties depend on the structure. For example, how it responds to pressure or how it can fold together. But Melio especially hopes that he can find a way to control the deformation of the microgrid remotely.

"Then you would actually have the basis for a microrobot. These are used, for example, in [biomedical applications](#), like operations. Of course, I'm not that far yet. I'm now experimenting with making the colloids magnetic to see if they can be controlled from the outside this way. It would be really nice if I could achieve that before I finish my Ph.D," says Melio.

**More information:** Julio Melio et al, Soft and Stiff Normal Modes in Floppy Colloidal Square Lattices, *Physical Review Letters* (2024). [DOI: 10.1103/PhysRevLett.132.078202](https://doi.org/10.1103/PhysRevLett.132.078202)

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