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Optical nanoresonators are fundamental building blocks in a number of nanotechnology applications (e.g. in spectroscopy) due to their ability to efficiently confine light at the nanoscale. Recently, nanoresonators based on the excitation of phonon polaritons (PhPs) – light coupled to lattice vibrations – in polar crystals (e.g. SiC or h-BN) have attracted much attention due to their strong field confinement, high-quality factors, and potential to enhance the photonic density of states at mid-infrared (IR) frequencies [1]. Here, we go one step further by introducing PhPs nanoresonators that not only exhibit these extraordinary properties but also incorporate a new degree of freedom: twist tuning, i.e. the possibility to be spectrally controlled by a simple rotation (Fig. 1a). To that end, we both take advantage of the low-loss-in-plane hyperbolic propagation of PhPs in the van der Waals crystal α -MoO₃ [2], and realize a dielectric engineering of a pristine α -MoO₃ slab placed on top of a metal ribbon grating, which preserves the high quality of the polaritonic resonances. By simply rotating the α -MoO₃ slab in the plane (from 0 to 45°), we demonstrate by far- and near-field measurements that the narrow polaritonic resonances (with quality factors Q up to 200) can be tuned in a broad range (up to 32cm⁻¹, i.e. up to~6 times its full width at half maximum, FWHM~5 cm⁻¹) (Fig 1b). Our results open the door to the development of tunable low-loss nanotechnologies at IR frequencies with application in sensing, emission, or photodetection.

References

Ma, W. et al., Nature, 562 (2018) 557-562. [2]

Figures



nanoresonators in α by placing a pristine

MoO₃ defined α -MoO₃ slab on top of metal ribbons. **a.** Schematics of the studied structure that allows defining the nanoresonators by ''dielectric engineering'' and controlling them by a twist angle, φ . **b.** Measured relative reflection spectra, δR for twist angles $\varphi = 0,15,30$ and 45° .

Figure 1: PhPs

Li, Peining. et al., Science, 359 (2018) 892-896. [1]