We fabricate artificial molecules composed of heavy atom lead on a van der Waals crystal. Pb atoms templated on a honeycomb charge-order superstructure of IrTe₂ form clusters ranging from dimers to heptamers including benzene-shaped ring hexamers. Tunneling spectroscopy and electronic structure calculations reveal the formation of unusual relativistic molecular orbitals within the clusters. The spin-orbit coupling is essential both in forming such Dirac electronic states and stabilizing the artificial molecules by reducing the adatom-substrate interaction. Lead atoms are found to be ideally suited for a maximized relativistic effect. This work initiates the use of novel two-dimensional orderings to guide the fabrication of artificial molecules of unprecedented properties.
Artificial atomic clusters or lattices supported on solid surfaces can exhibit tailored electronic, magnetic, and topological properties of fundamental and technological importance. For example, atomic-scale chains and clusters disclosed the quantum confinement of electrons, topological edge modes, superlattice Dirac bands, flat bands, atomic-scale spin interactions, and topological defects. The direct atom-by-atom manipulation and the self-assembly of atoms or molecules are two major approaches to realize such atomic-scale chains, clusters, and finite lattices. Both methods, however, have their own limitations under given interatomic interactions, and fabricating energetically and kinetically unfavorable structures or assemblies has been a huge challenge.

In case of the self-assembly of supported clusters, such limitations may be overcome by templates, which provide unusual growth environments to produce otherwise unfavored molecular structures. Step edges and one-dimensional surface superstructures were actively used as such templates on metal and Si surfaces. The growth of metal atoms on these substrates leads to unusually anisotropic clusters. On the other hand, nanosized two-dimensional superstructures were suggested as novel templates, such as the moiré structure of hexagonal boron nitride, metal–organic frameworks of DNA, and metal–organic honeycomb networks on metal surfaces.

However, while the confinement of metal atoms and molecules on these templates were demonstrated, the formation of artificial molecules was not reported yet.

In this work, we employ a novel template-guided atomic self-assembly technique and take advantage of the strong relativistic effects in lead atoms that effectively create long-ranged interatomic bonds. This allows us to produce a series of unprecedented artificial molecules with relativistic, so-called Dirac, electronic orbitals. The honeycomb charge-order superstructure of the van der Waals (vdW) crystal IrTe$_2$ cages various Pb clusters ranging from dimers to heptamers, including those in interesting benzene-like hexagonal rings. Atomically resolved spectroscopy and electronic structure calculations reveal clearly the formation of molecular orbitals by direct orbital overlap between neighboring Pb adatoms. The spin–orbit coupling (SOC) of Pb atoms and of the substrate is shown to enhance the interatomic interaction to drive the formation of unprecedentedly relativistic molecules at an unusually large interatomic distance. This case exemplifies the chemical stabilization of nanoscale assemblies uniquely by relativistic effects and opens the potential of novel superstructures on vdW crystals for the fabrication of otherwise unfavored molecules with unprecedented properties.

**Results**

**Charge-order superstructure hosts Pb clusters.** Figure 1a shows Pb adatoms deposited on the honeycomb charge-order superstructure at the surface of a IrTe$_2$ vdW crystal. Below 180 K, a honeycomb superstructure forms with a period of 2.2 nm by the spontaneous ordering of 5$d^{3+}$ and 5$d^{4+}$ valence electrons of Ir (with the buckling of the Te layer accompanied) and competes with various stripe charge orders. Pb adatoms sit at hollow sites surrounded by three Te atoms (Fig. 1b, Supplementary Fig. 2) and, very selectively with yet unclear reasons, inside of each honeycomb, which are above Ir atoms with the 5$d^{4+}$ valency (Supplementary Fig. 1). Density-functional theory (DFT) calculations, including SOC consistently predict the four-fold hollow site adsorption with an adsorption energy of $-0.93$ eV against the three-fold hollow site of $-0.74$ eV and Te on-top site of $-0.16$ eV. Clusters ranging from one to seven Pb atoms are confined within a honeycomb unitcell with various lateral configurations. The Pb–Pb interatomic distance is dictated by the substrate with two choices of $\sqrt{3}a_0$ or $2a_0$ ($a_0$, the IrTe$_2$ lattice constant of 3.5 Å (ref. 26)). The $2a_0$-spaced dimers (Fig. 1c) are more frequently observed than those of $\sqrt{3}a_0$ and Pb–Pb distance (Fig. 1d). As trimers, a heterogeneous trimer with a Pb–Pb spacing both of $\sqrt{3}a_0$ and $2a_0$, or a compact one of $2a_0$ or $\sqrt{3}a_0$ are formed (Fig. 1e, Supplementary Fig. 4c). Figure 1f, g shows a tetramer formed by two $2a_0$ dimers, and one with $\sqrt{3}a_0$ and $2a_0$ dimers, respectively. We also find benzene-like hexagonal ring clusters as shown in Fig. 1h–j among a few other interesting clusters. These configurations are made only by $2a_0$ building blocks and are the largest clusters confined within the hexagonal unit of the template.

**Relativistic molecular orbitals in Pb dimers.** Electronic energy levels of Pb clusters are revealed by scanning tunneling spectroscopy (STS). For a monomer (Fig. 2c), we observe three main spectral features at 1.40, 1.71, and 1.93 eV for Pb 6$p$ valence electrons. Our calculation shows that a Pb adatom is partly ionized by donating 6$p$ electrons into the substrate to shift the 6$p$ states to unoccupied states (Fig. 2o, w). As shown in Fig. 2w, the
relativistic effect of SOC splits the $6p$ states into $6p_{1/2}$ and $6p_{3/2}$ Dirac orbitals and the latter splits further due to the substrate-induced splitting of in-plane ($p_{xy}$) and out-of-plane ($p_z$) orbitals. Consequently, three main peaks correspond to the relativistic $p$ orbitals of $p_{1/2}$ ($j = 1/2, m_j = \pm 1/2$) and $p_{3/2}$ ($j = 3/2, m_j = \pm 3/2, \pm 1/2$), respectively.

In case of a dimer with a shorter Pb–Pb distance, an energy splitting ($0.23$ eV) of molecular bonding and antibonding states is observed clearly for the $6p_{1/2}$ state (Fig. 2). In a longer dimer, the splitting is much smaller, $\approx0.10$ eV, as the orbital overlap reduces (Fig. 2f). Our DFT calculation predicts (Fig. 2r, v) the lowest molecular orbital splittings of $0.05$ and $0.18$ eV for the longer and shorter dimers, respectively, in reasonable agreement with the experiment. The present DFT calculation seems to underestimate the molecular bonding interaction by $\approx0.05$ eV. These energy scale is consistent with what are expected for a freestanding Pb–Pb dimer (Supplementary Fig. 6). This result clearly indicates the molecular orbital formation within the Pb clusters at an unusually large Pb–Pb distance of $7\,\text{Å}$, which is observed consistently for all cluster configurations identified (Supplementary Fig. 7).

The molecular levels of a Pb dimer are well explained by the simple tight-binding interaction of Dirac atomic orbitals perturbed weakly by the substrate as discussed above. That is, three relativistic $p$ orbitals split simply into the combination of molecular orbitals in gerade ($g$) and ungerade ($u$) symmetries (Fig. 2w). For example, the two lowest peaks correspond to the combination of the $p_{1/2}$ orbitals in $g$ and $u$ symmetries. The resulting Dirac molecular orbitals are clearly differentiated from their scalar-relativistic counterparts of $\sigma$ and $\pi$ molecular orbitals. The $g$ ($u$) orbital consists of one-third $\sigma$-bond ($\sigma$-antibond) and two-thirds $\pi$-bond ($\pi$-antibond) character, indicating a substantial mixing between $\sigma$- and $\pi$-bonds due to the full relativity effect of SOC. The Dirac basis and the interaction with substrate atoms also breaks the mirror symmetry perpendicular to the dimer axis. This is reasonably confirmed in the experiment (Fig. 2h, i) and in the calculation (Fig. 2t, u). Further details of the interaction with the substrate will be discussed below, which become important for the understanding of the relativistic effect on the adsorption energy.

The direct and real space observation of relativistic molecular orbitals is exceptional, since most of previous experiments for
Mechanism of the interaction between Pb atoms. It is important to point out that the attractive interaction between Pb adatoms is only due to SOC and that the 2a₀ distance imposed by the substrate is optimal for the formation of a Pb dimer (Fig. 2x) in our calculations. This is consistent with the observation of the higher population of 2a₀ dimers. The SOC energetics for the larger Pb molecules, from trimers to heptamer (Fig. 2y, Supplementary Fig. 10), is also consistent with the experimental observation of larger Pb molecules with exclusively 2a₀ nearest-neighbor distance. The 2a₀ distance is optimized by two competing energy contributions, the ionic (or dipole) repulsion and the orbital overlap energy gain of adatom–substrate hybridized states. The comparison of the calculated differential charges with and without SOC reveals two sophisticated SOC effects for those contributions: (i) a reduced donation of p electrons to the substrate to reduce the ionic repulsion and (ii) an enhanced orbital overlap of the Pb–Te hybridized states (Supplementary Fig. 10). The former is due to the lowering of the atomic energy level of Dirac orbitals by SOC. It is also partly due to the charge redistribution of the IrTe₂ substrate by SOC, namely the charge transfer from the dₐ states of Ir atom to Te atoms. The substantial interaction of Pb with the substrate Te orbitals provides an extra energy gain for the antibonding level of the p₁/₂ (mₗ = ±3/2) orbital of the 3a₀ dimer (Supplementary Fig. 9b). That is, the interwined effect of the substrate and the relativity determines the stability of the Pb molecules. In these respects, Pb is ideal since it has very strong relativistic effects (in comparison, for example, to Sn, Supplementary Fig. 11) and a proper interaction strength with the substrate (in comparison to Tl, Supplementary Fig. 12). Of course, the formation of Pb molecules would also be affected by the size and shape of the charge-order honeycomb template.

Benzenelike ring-shaped Dirac molecules. While a consistent interaction mechanism holds for all 15 different Pb molecules identified (Supplementary Fig. 4), particularly interesting ones are those based on a six-fold hexagonal ring (Fig. 3). The benzene-like ring molecule of Pb₉ with a 2a₀ Pb–Pb distance exhibit the weak molecular splitting similar to a 2a₀ dimer (Fig. 3e). The intriguing molecular orbital formation is clearly imaged by the dill/DV maps: (i) the broken AB sublattice symmetry for the lowest-energy state (Fig. 3b), (ii) a Kekulé-like distortion for the second state (Fig. 3c), and (iii) a mirror symmetry-broken two-fold symmetric bonding feature for the third state (Fig. 3d). Our calculation reproduces well the spatial characteristics of these molecular orbitals, which are indeed due to the relativistic character and the interaction with the substrate (Fig. 3l–n, Supplementary Fig. 13). Note, however, the remaining energetic discrepancy between experiment and calculation. The present calculations underestimate the molecular bonding interaction and the calculated spectral features are almost degenerate at ~1.3 eV for the benzene-like molecule. We think that this difference is at least partly due to the limitation of our model in taking into account of the substrate electronic states in its charge-ordered correlated state. Another interesting molecule is the Pb pentamer (Fig. 3f–j), whose Dirac molecular orbitals are also reasonably well simulated (Fig. 3p–t). Such type of an individual open ring molecule cannot be synthesized. The Pb₂ molecule features the edge states at both truncated ends between the bonding and antibonding levels of three Dirac p orbitals (Supplementary Fig. 14). It indicates the ring-type Pb molecules are circularly delocalized electron systems. The formation of a filled-benzene-ring molecule is also highly unusual (Fig. 1j), where the interatomic overlap is much enhanced by the central Pb atom with unique bonding configurations (Supplementary Figs. 15–17). This molecule can be compared with a rare example of a planar B₇ molecule except for the strong SOC. While not accessible by the present experiments, the benzene-type Dirac molecular orbitals fabricated here have unique and interesting spin configurations (Supplementary Fig. 13), which may be exploited further. It becomes very obvious that the present artificial Pb molecules introduce unprecedented molecular configurations combined with the strong SOC.

Summary. We fabricate various artificial Pb molecules utilizing a novel honeycomb template of the charge-ordered superstructure in IrTe₂. The molecular orbital formation is clearly observed in the Pb clusters from dimers to unusual benzene-ring-type molecules. The SOC is found to play crucial roles in both the condensation of Pb atoms and the molecular orbital formation, bringing them into an unprecedented regime of relativistic Dirac molecules. Beyond the relativistic chemistry, the unusual bonding and spin structure of these relativistic molecular orbitals may be combined with the novel electronic properties of the substrate, such as the charge ordering and the emerging superconductivity to lead to a new type of a quantum system. In addition to the versatile technique of atom-by-atom manipulation, where the interatomic distance may be varied, the use of various types of 2D superstructures as templates would definitely extend the potential of molecular or cluster self-assemblies. Interesting 2D superstructures can include twisted bilayer graphene and domain-wall or twin-boundary networks of transition metal dichalcogenides. The templated self-assembly of magnetic atoms on such substrates can be very interesting due to the proximity to the novel 2D electronic states of substrates.

Methods
Sample preparation. Single crystals of IrTe₂ were grown by Te flux using presintered IrTe₂ polycrystals, as reported previously. Samples were cleaved in a vacuum better than 5 × 10⁻¹⁰ torr at room temperature. Pb atoms are grown on cleaved IrTe₂ surfaces at room temperature. The Pb atom tends to cover the IrTe₂ surface up to 0.1 mL. Over this coverage, this Pb growth mode depends on the deposition temperature. For example, when the Pb grows at room temperature, Pb forms a uniform film. However, at low-temperature deposition, depending on the coverage and annealing condition, we can find different heights of the Pb islands.

STM measurement. All the STM measurements were performed with a commercial ultrahigh vacuum cryogenic STM (Specs, Germany) in the constant-current mode with PtIr tips at 4.3 K. The differential conductance, dI/dV, was measured using the lock-in detection with a modulation of 1.17 kHz.

DFT calculations. We perform the relativistic DFT calculations using the Vienna abinitio simulation package within the generalized gradient approximation of the Perdew–Burke–Ernzerhof type and the projector augmented-wave method. The IrTe₂ surface is modeled by a (5 × 5) and (7 × 7) supercell with a single IrTe₂ layer and a vacuum spacing of ~23.6 Å. The calculated value 3.839 Å is used as the lattice constant of IrTe₂. We use a plane-wave basis with a cutoff of 211 eV for expansion of electronic wave functions and a 15 × 15 × 1 k-point mesh for the 1 x 1 Brillouinzone integrations. To avoid the unexpected substrate distortion of the IrTe₂, only adsorbed Pb atoms are relaxed until the residual force components are within 0.02 eVÅ. Here, the Fermi level was shifted down by 0.70 eV because to overcome the inherent discrepancy between experiment and theory due to unknown charge-ordered structure, doping effect of the Pb adsorbates and the IrTe₂ sample and/or the inaccuracy of DFT band-energy calculations.

very few heavy element molecules relied on spectroscopy techniques and, thus, the relativistic effects were mainly discussed in theoretical aspects. Although a Pb dimer has a much longer bond length (7 Å) than other relativistic molecules and there is a substantial adatom–substrate interaction, the molecular splitting mainly arises from the direct overlap of relativistic p orbitals. This is distinguished from the substrate-mediated long-range interaction of other metal clusters on metal surfaces.
given energies (spin decoupled and spin summed, respectively), and

6. Stroscio, J. A. et al. Electronically induced atom motion in engineered CoCu

and black Pb atoms indicated in

4. Wallis, T. M., Nilius, N., Mokaelian, G. & Ho., W. Electronically induced atom motion in engineered CoCu, respectively.

5. Lagoute, J., Liu, X. & Fölsch, S. Link between adatom resonances and the Cu states at truncated end atoms (the other Pb atoms). Solid and dashed lines represent

For the Pb hexamer, the localized DOS at red and blue Pb atoms is identical to each other. For the Pb pentamer, the red (dark blue) line denote the localized states at truncated end atoms (the other Pb atoms).

Data availability
The authors declare that the data supporting the findings of this study are available within the article and its Supplementary Information.

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References
1. Nilius, N., Wallis, T. M. & Ho., W. Development of one-dimensional band structure in artificial gold chains. Science 297, 1853 (2002).
2. Nilius, N., Wallis, T. M., Person, M. & Ho., W. Distance dependence of the interaction between single atoms: gold dimers on NiAl(110). Phys. Rev. Lett. 90, 196103 (2003).
3. Fölsch, S., Hyldgaard, P., Koch, R. & Ploog, K. H. Quantum confinement in monatomic Cu chains on Cu(111). Phys. Rev. Lett. 92, 56803 (2004).
4. Wallis, T. M., Nilius, N., Mokaelian, G. & Ho., W. Electronic properties of artificial Au chains with individual Pd impurities. J. Chem. Phys. 122, 011101 (2005).
5. Lagoute, J., Liu, X. & Fölsch, S. Link between adatom resonances and the Cu (111) shockley surface state. Phys. Rev. Lett. 95, 136801 (2005).
6. Stroscio, J. A. et al. Electronically induced atom motion in engineered CoCu nanostructures. Science 313, 948 (2006).
7. Nadj-Perge, S. et al. Observation of Majorana fermions in ferromagnetic atomic chains on a superconductor. Science 346, 602 (2014).
8. Yankowitz, M. et al. Emergence of superlattice Dirac points in graphene on hexagonal boron nitride. Nat. Phys. 8, 382 (2012).
9. Slot, M. R. et al. Experimental realization and characterization of an electronic Lieb lattice. Nat. Phys. 13, 372 (2017).
10. Hirjibehedin, C. F., Christopher, P. L. & Heinrich, A. J. Spin coupling in engineered atomic structures. Science 312, 1021 (2006).
11. Cheon, S., Kim, T.-H., Lee, S.-H. & Yeom, H. W. Chiral solitons in a coupled double Peierls chain. Science 350, 182 (2015).
12. Drost, R., Ojanen, T., Harju, A. & Liljeroth, P. Topological states in engineered atomic lattices. Nat. Phys. 13, 668 (2017).
13. Chichak, K. S. et al. Molecular Borromean rings. Science 304, 1308 (2004).
14. Lobo, J. et al. Tuning the surface state dimensionality of Cu nanostripes. Phys. Rev. Lett. 93, 137602 (2004).
15. Hermes, S., Schröder, F., Chelmowski, R., Woll, C. & Fischer, R. A. Selective nucleation and growth of metal-organic open framework thin films on patterned COOH/CF3- terminated self-assembled monolayers on Au(111). J. Am. Chem. Soc. 127, 13744 (2005).
16. Barth, J. V., Costantini, G. & Kern, K. Engineering atomic and molecular nanostructures at surface. Nature 437, 671 (2005).
17. Kang, P.-G., Jeong, H. & Yeom., H. W. Microscopic mechanism of templated self-assembly: Indium metallic atomic wires on Si(553)-Au. Phys. Rev. B 79, 113403 (2009).
18. Corso, M. et al. Boron nitride nanomesh. Science 303, 217 (2004).
19. Dilk, H. et al. Surface trapping of atoms and molecules with dipole rings. Science 319, 1824 (2008).
20. LaBean, T. H. et al. DNA-Templated self-assembly of protein arrays and highly conductive nanowires. Science 301, 5641 (2003).
24. Nowakowska, S. et al. Interplay of weak interactions in the atom-by-atom
   adsorption process. Phys. Rev. Lett. 105, 066104 (2010).
23. Pivetta, M. et al. Formation of Fe cluster superlattice in a metal-organic
   molecular network on metal surface. J. Phys. Condens. Matter 28, 153003 (2016).
22. Cheng, Z. et al. Adsorbates in a box: titration of substrate electronic states.
   Phys. Rev. Lett. 110, 086102 (2013).
21. Kiehl, R. A. et al. DNA-Templated self-assembly of metallic nanocomponent
   arrays on a surface. Nano Lett. 4, 2343 (2004).
20. Knorr, N. et al. Effect of spin-orbit coupling on the electronic
   structure and excitations of the Bi− anion in (K-crypt)2 Bi, on the basis
   of relativistic electronic structure calculations. J. Phys. Chem. A 109, 1675 (2005).
19. van Lenthe, E., Snijders, J. G. & Baerends, E. J. The zero-order regular
   approximation for relativistic effects: the effect of spin-orbit coupling in closed
   shell molecules. J. Chem. Phys. 105, 6505 (1996).
18. Iliaš, M., Kelló & Urban, M. Relativistic effects in atomic and molecular
   properties. Acta Physica Slovaca 60, 259 (2010).
17. Alexandrova, A. N., Boldyrev, A. I., Zhai, H.-J. & Wang, L.-S. Electronic
   structure, isomerism, and chemical bonding in Bi, and CH2 BrCN by collision with He
   metastable atoms. J. Phys. Chem. A 107, 2137 (2003).
16. Daï, D. et al. Analysis of the effect of spin-orbit coupling on the electronic
   structure and excitation spectrum of the Bi− anion in (K-cript)2 Bi, on the basis
   of relativistic electronic structure calculations. J. Phys. Chem. A 109, 1675 (2005).
15. Gans, B., Grassi, G. & Merkt, F. Spin-orbit and vibronic coupling in the ionic
   ground state of iodoacetylene from a rotationally resolved photoelectron
   spectrum. J. Phys. Chem. A 117, 9353 (2012).
14. Iliaš, M. Relativistic effects in atomic and molecular
   properties. Acta Physica Slovaca 60, 259 (2010).
13. van Lenthe, E., Snijders, J. G. & Baerends, E. J. The zero-order regular
   approximation for relativistic effects: the effect of spin-orbit coupling in closed
   shell molecules. J. Chem. Phys. 105, 6505 (1996).
12. Huda, M. N., Niranjan, M. K., Sahu, B. R. & Klenman, L. Effect of spin-orbit
   coupling on small platinum nanoclusters. Phys. Rev. A 73, 053201 (2006).
11. Li, W.-L., Lu, J.-B., Wang, Z.-L., Hu, H.-S. & Li, J. Relativity-induced bonding
   pattern change in coinage metal dimers M2 (M = Cu, Ag, Au, Rg). Inorg.
   Chem. 57, 5499 (2018).
10. Repp, J., MoreSCO, F., Meyer, G. & Rieder, K.-H. Substrate mediated long-range
   oscillatory interaction between adatoms: Cu/Cu(111). Phys. Rev. Lett. 85, 2981 (2000).
9. Knorr, N. et al. Long-range adsorbate interactions mediated by a two-
   dimensional electron gas. Phys. Rev. B 65, 115420 (2002).
8. Alexandrova, A. N., Boldyrev, A. I., Zhai, H.-J. & Wang, L.-S. Electronic
   structure, isomerism, and chemical bonding in Bi, and Re, J. Phys. Chem. A 108, 3509 (2004).
7. Mele, E. J. Commensuration and interlayer coherence in twisted bilayer
   graphene. Phys. Rev. B 81, 161405(R) (2010).
6. Cho, D. et al. Nanoscale manipulation of the Mott insulating state coupled to
   charge order in 1T-TaS2. Nat. Commun. 7, 10453 (2016).
5. Ma, Y. et al. Metallic twin grain boundaries embedded in MoSe2 monolayers
   grown by molecular beam epitaxy. ACS Nano 11, 5130 (2017).
4. Kresse, G. & Furthmüller, J. Efficient iterative schemes for ab initio total
   energy calculations using a plane-wave basis set. Phys. Rev. B 54, 11169 (1996).
3. Perdew, J. P., Burke, K. & Ernzerhof, M. Generalized gradient approximation
   made simple. Phys. Rev. Lett. 77, 3865 (1996).
2. Blochl, P. E. Projector augmented-wave method. Phys. Rev. B 50, 17953 (1994).

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Author contributions
J.W.P. performed the DFT calculations. H.S.K. carried out the STM measurements. J.W. P. and H.W.Y. analyzed the data, and wrote the manuscript with the comments of all other authors. H.W.Y. supervised the research.

Competing interests
The authors declare no competing interests.

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