



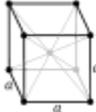
decay;<sup>[21]</sup> however, no known roentgenium isotope has been observed to undergo beta decay.<sup>[17]</sup> The unknown isotopes <sup>277</sup>Rg and <sup>283</sup>Rg are also expected to have long half-lives of 1 second and 10 minutes respectively. Before their discovery, the isotopes <sup>278</sup>Rg, <sup>281</sup>Rg, and <sup>282</sup>Rg were predicted to have long half-lives of 1 second, 1 minute, and 4 minutes respectively; however, they were discovered to have shorter half-lives of 4.2 milliseconds, 17 seconds, and 2.1 minutes respectively.<sup>[17]</sup>

## Predicted properties

### Chemical

Roentgenium is the ninth member of the 6d series of transition metals. Since copernicium (element 112) has been shown to be a transition metal, it is expected that all the elements from 104 to 112 would form a fourth transition metal series.<sup>[22]</sup> Calculations on its ionization potentials and atomic and ionic radii are similar to that of its lighter homologue gold, thus implying that roentgenium's basic properties will resemble those of the other group 11 elements, copper, silver, and gold; however, it is also predicted to show several differences from its lighter homologues.<sup>[2]</sup>

Roentgenium is predicted to be a noble metal. Based on the most stable oxidation states of the lighter group 11 elements, roentgenium is predicted to show stable +5, +3, and −1 oxidation states, with a less stable +1 state. The +3 state is predicted to be the most stable. Roentgenium(III) is expected to be of comparable reactivity to gold(III), but should be more stable and form a larger variety of compounds. Gold also forms a somewhat stable −1 state due to relativistic effects, and roentgenium may do so as well:<sup>[2]</sup> the electron affinity of roentgenium is expected to be around 1.6 eV (37 kcal/mol), significantly lower than gold's value of 2.3 eV (53 kcal/mol), so roentgenides may not be stable or even possible.<sup>[4]</sup> The 6d orbitals are destabilized by relativistic effects and spin-orbit interactions near the end of the fourth transition metal series, thus making the high oxidation state roentgenium(V) more stable than its lighter homologue gold(V) (known only in one compound) as the 6d electrons participate in bonding to a greater extent. The spin-orbit interactions stabilize

|  | 3rd: 3077.9 kJ/mol (more) <i>(all estimated)</i> <sup>[2]</sup>   |   |          |          |                   |
|--|---|---|----------|----------|-------------------|
| <b>Atomic radius</b>                       | empirical: 138 pm <i>(predicted)</i> <sup>[2][4]</sup>  |   |          |          |                   |
| <b>Covalent radius</b>                     | 121 pm <i>(estimated)</i> <sup>[5]</sup>  |   |          |          |                   |
| <b>Miscellanea</b>                         |   |   |          |          |                   |
| <b>Crystal structure</b>                   | body-centered cubic (bcc) <i>(predicted)</i> <sup>[3]</sup>  |   |          |          |                   |
| <b>CAS Number</b>                          | 54386-24-2  |   |          |          |                   |
| <b>History</b>                             |   |   |          |          |                   |
| <b>Naming</b>                              | after Wilhelm Röntgen   |   |          |          |                   |
| <b>Discovery</b>                           | Gesellschaft für Schwerionenforschung (1994)  |   |          |          |                   |
| <b>Most stable isotopes of roentgenium</b> |   |   |          |          |                   |
| iso  | NA  | half-life                               | DM       | DE (MeV) | DP                |
| <sup>282</sup> Rg <sup>[6]</sup>           | syn   | 2.1 <sup>+1.4</sup> <sub>−0.6</sub> min | α        | 9.00     | <sup>278</sup> Mt |
| <sup>281</sup> Rg <sup>[7][8]</sup>        | syn   | 17 <sup>+6</sup> <sub>−3</sub> s        | SF (90%) |          |                   |
|  |   |   | α (10%)  |          | <sup>277</sup> Mt |
| <sup>280</sup> Rg                          | syn   | 3.6 s                                   | α        | 9.75     | <sup>276</sup> Mt |
| <sup>279</sup> Rg                          | syn   | 0.17 s                                  | α        | 10.37    | <sup>275</sup> Mt |

molecular roentgenium compounds with more bonding 6d electrons; for example,  $\text{RgF}_6^-$  is expected to be more stable than  $\text{RgF}_4^-$ , which is expected to be more stable than  $\text{RgF}_2^-$ . Roentgenium(I) is expected to be difficult to obtain.<sup>[2][23][24]</sup>

The probable chemistry of roentgenium has received more interest than that of the two previous elements, meitnerium and darmstadtium, as the valence s-subshells of the group 11 elements are expected to be relativistically contracted most strongly at roentgenium.<sup>[2]</sup> Calculations on the molecular compound  $\text{RgH}$  show that relativistic effects double the strength of the roentgenium–hydrogen bond, even though spin-orbit interactions also weaken it by 0.7 eV (16 kcal/mol). The compounds  $\text{AuX}$  and  $\text{RgX}$ , where  $X = \text{F, Cl, Br, O, Au, or Rg}$ , were also studied.<sup>[2][25]</sup>  $\text{Rg}^+$  is predicted to be the softest metal ion, even softer than  $\text{Au}^+$ , although there is disagreement on whether it would behave as an acid or a base.<sup>[26][27]</sup> In aqueous solution,  $\text{Rg}^+$  would form the aqua ion  $[\text{Rg}(\text{H}_2\text{O})_2]^+$ , with an Rg–O bond distance of 207.1 pm. It is also expected to form  $\text{Rg}(\text{I})$  complexes with ammonia, phosphine, and hydrogen sulfide.<sup>[27]</sup>

## Physical and atomic

Roentgenium is expected to be a solid under normal conditions and to crystallize in the body-centered cubic structure, unlike its lighter congeners which crystallize in the face-centered cubic structure, due to its being expected to have different electron charge densities from them.<sup>[3]</sup> It should be a very heavy metal with a density of around  $28.7 \text{ g/cm}^3$ ; in comparison, the densest known element that has had its density measured, osmium, has a density of only  $22.61 \text{ g/cm}^3$ . This results from roentgenium's high atomic weight, the lanthanide and actinide contractions, and relativistic effects, although production of enough roentgenium to measure this quantity would be impractical, and the sample would quickly decay.<sup>[2]</sup>

The stable group 11 elements, copper, silver, and gold, all have an outer electron configuration  $nd^{10}(n+1)s^1$ . For each of these elements, the first excited state of their atoms has a configuration  $nd^9(n+1)s^2$ . Due to spin-orbit coupling between the d electrons, this state is split into a pair of energy levels. For copper, the difference in energy between the ground state and lowest excited state causes the metal to appear reddish. For silver, the energy gap widens and it becomes silvery. However, as the atomic number increases, the excited levels are stabilized by relativistic effects and in gold the energy gap decreases again and it appears gold. For roentgenium, calculations indicate that the  $6d^97s^2$  level is stabilized to such an extent that it becomes the ground state and the  $6d^{10}7s^1$  level becomes the first excited state. The resulting energy difference between the new ground state and the first excited state is similar to that of silver and roentgenium is expected to be silvery in appearance.<sup>[1]</sup> The atomic radius of roentgenium is expected to be around 138 pm.<sup>[2]</sup>

## Experimental chemistry

Unambiguous determination of the chemical characteristics of roentgenium has yet to have been established<sup>[28]</sup> due to the low yields of reactions that produce roentgenium isotopes.<sup>[2]</sup> For chemical studies to be carried out on a transactinide, at least four atoms must be produced, the half-life of the isotope used must be at least 1 second, and the rate of production must be at least one atom per week.<sup>[22]</sup> Even though the half-life of <sup>281</sup>Rg, the most stable known roentgenium isotope, is 26 seconds, long enough to perform chemical studies, another obstacle is the need to increase the rate of production of roentgenium isotopes and allow experiments to carry on for weeks or months so that statistically significant results can be obtained. Separation and detection must be carried out continuously to separate out the roentgenium isotopes and automated systems can then experiment on the gas-phase and solution chemistry of roentgenium as the yields for heavier elements are predicted to be smaller than those for lighter elements. However, the experimental chemistry of roentgenium has not received as much attention as that of the heavier elements from copernicium to livermorium,<sup>[2][28][29]</sup> despite early interest in theoretical predictions due to the maximizing of relativistic effects on the *ns* subshell in group 11 occurring at roentgenium.<sup>[2]</sup>

## Source

- Wikipedia: Roentgenium (<https://en.wikipedia.org/wiki/Roentgenium>)