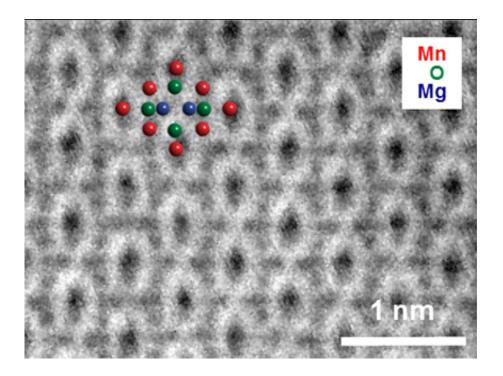


## "High-occupancy" vehicle (HOV) battery

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The location of magnesium inserted into a battery electrode material is shown in an atomic-resolution scanning transmission electron microscopy (STEM) image. The characteristic "diamond" unit cell is overlaid, indicating the atomic positions of manganese (red), oxygen (green), and magnesium (blue); clear occupancy of specific sites within the crystal, called tetrahedral sites, is shown. Credit: Chunjoong Kim, University of Illinois

In lithium-ion batteries, a singly charged lithium atom (or "ion") flows between two electrodes while one electron is contributed to the electrical current in the external circuit. If too many lithium ions are intercalated (inserted between existing atomic layers), the electrode's atomic



structure can distort, which usually limits the ability to recharge the battery. In a battery electrode composed of manganese oxide  $(Mn_2O_4)$ , magnesium ions with two positive charges  $(Mg^{2+})$  could provide two electrons per intercalated ion—twice the electrical current if present at the same density—without irreversible distortion of the electrode material. This research showed that  $Mg^{2+}$  intercalation into  $Mn_2O_4$  was extensive and reversible.

Future battery systems composed of cathodes (negatively charged electrodes) that take advantage of  $Mg^{2+}$  intercalation could have twice or more the energy density of today's commercial <u>lithium-ion</u> batteries. This new class of batteries may lead to longer-lasting cell phones and more practical electric vehicles.

Batteries are composed of two electrodes, an anode (positive charge) and a cathode (negative charge), separated by an electrical insulator. As the battery operates with electrical current flowing in the external circuit, positively charged atoms (or ions) flow internally between the two electrodes. Intercalation of too many ions from the electrolyte into the electrodes can cause distortion, which can limit the ability to recharge the battery. Batteries using Mg<sup>2+</sup> ions instead of singly charged lithium ions have the potential for significantly increased energy density and the production of two electrons of electrical current for each intercalated ion. A major challenge for magnesium batteries is finding suitable cathode materials capable of storing a large amount of Mg<sup>2+</sup> ions while operating at high voltage. This research demonstrates that Mn<sub>2</sub>O<sub>4</sub> with the spinel structure is such a material. Researchers at the Joint Center for Energy Storage Research (JCESR) found that Mg intercalation into the electrode was extensive and reversible. The scientists used four techniques to characterize this phenomenon at multiple length and chemical scales. Electron microscopy with elemental analysis permitted visualization of the electrochemical intercalation. X-ray diffraction allowed confirmation of the expected structural transition upon



intercalation. Nuclear magnetic resonance (NMR) and X-ray absorption spectroscopy (XAS) further validated the Mg intercalation. This research has confirmed the validity of a new class of materials for battery cathodes based on  $Mg^{2+}$  intercalation. Coupled with a pure Mg anode, these battery systems could have twice or more the <u>energy density</u> of today's commercial lithium-ion batteries along with increased safety and reduced cost.

**More information:** Chunjoong Kim et al. Direct Observation of Reversible Magnesium Ion Intercalation into a Spinel Oxide Host, *Advanced Materials* (2015). DOI: 10.1002/adma.201500083

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