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Cs-corrected STEM Observation for Al-Rh-Cu Decagonal Quasicrystal

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On the basis of Cs-corrected high-angle annular detector dark-field (HAADF)- and annular bright-field (ABF)scanning transmission electron microscopy (STEM) observation, the structure of an Al-Rh-Cu decagonal quasicrystal (DQC) formed with two quasiperiodic planes along the periodic axis in an annealed $Al_{63}Rh_{18,5}Cu_{18,5}$ alloy has been investigated. Heavy atoms of Rh, and mixed sites (MSs) of Al and Cu atoms projected along the periodic axis can be clearly recognized as separate bright dots in HAADF-STEM images, and consequently arrangements of Rh atoms and MSs on the two quasiperiodic planes can be directly determined from those of bright dots in HAADF-STEM images. The Rh atoms are arranged in pentagonal tiling formed with pentagonal and star-shaped pentagonal tiles with an edge-length of 0.76 nm, and also MSs with a pentagonal arrangement are located in the pentagonal tiles with definite orientations. The star-shaped pentagonal tiles in the pentagonal tiling are arranged in $\tau^2(\tau;$ golden ratio)-inflated pentagonal tiling with a bond-length of 2 nm. From arrangements of Rh atoms located in pentagonal tilings with a bond-length of 2 nm, which are generated by the projection of a hyper-cubic lattice in a five-dimensional superspace, occupation domains in the perpendicular space are derived. Al atoms as well as Rh atoms and MSs are represented as dark dots in ABF-STEM images. The present model of the Al-Rh-Cu DQC is basically different from the structure proposed by single crystal Xray diffraction in respect of the presence of the 0.76 nm pentagonal tiling of Rh atoms and MSs of Al and Cu atoms.

Mo-PO8

The crystal structure of two novel cage-like aluminides *

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Two novel intermetallic compounds from Al-rich region of the Ce-Rh-Al phase diagram were synthesized using arc-melting technique. The crystal structure of Ce₂Rh₃Al₁₄ was analyzed by means of powder X-ray diffraction and Rietveld analysis. In the case of Ce₃Rh₇Al₂₉, structure determination was performed using single-crystal X-ray diffraction data. Single crystals were selected from crashed polycrystalline ingots.

The compound Ce₃Rh₇Al₂₉ of the new structure type crystallizes in orthorhombic space group (SG) *Cmcm*, Pearson symbol oC156 with a=9.1570(4), b=31.3790(14), c=8.9414 (17) Å, V=2569.2(17) Å³. Intermetallics Ce₂Rh₃Al₁₄ has a tetragonal structure closely related to Ce₂Ru₃Al₁₅[†], SG *I*4₁/*amd*, Pearson symbol *tI*136, a=9.0000(15), c=31.7640(11) Å), V=2572.9(6) Å³. Three cerium atoms (4c site) in Ce₃Rh₇Al₂₉ and two cerium atom (8e site) in Ce₂Rh₃Al₁₄ are coordinated by 18 and 20 Rh and Al atoms, forming the 3D- frameworks of large icosahedral voids. This feature resembles the high coordinated Ce environment seen in CeRh₄Al_{15.7} and CeRh₂Al₁₀. With the exception of the disordered Ce1 in Ce₃Rh₇Al₂₉, the closest Ce-Ce contacts in both compounds are about 4.5 Å.

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