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the x-ray diffraction pattern of anaphtite was used to model the new material. the starting model was an orthorhombic unit cell. the refined parameters are as follows: unit cell constants $a=6.5471(10)\text{\AA}$, $b=8.2459(12)\text{\AA}$, $c=3.6917(5)\text{\AA}$; the number of formula units per unit cell is 1. the crystal structure of the new product is monoclinic and belongs to the space group $p21/n$. a detailed structure analysis of the new product has been performed using rietveld refinement, single crystal x-ray diffraction and fullprof software. the crystal structure of the new product has been solved from the powder x-ray diffraction data by simulated annealing of eight independent structural sub-units, revealing a combination of four- and five-fold coordinated aluminums within the crystal structure. the software package is basically an x-ray diffraction data analysis program for determining the crystal structure, phase content, crystallite size, lattice parameters, and other information of the material and the accuracy of the analysis is dependent on the quality of the data and the selected instrumental configuration. this article describes the methodology for the determination of the structure of the materials and the accuracy of the analysis is dependent on the quality of the data and the selected instrumental configuration. the software package is essentially a high-throughput tool for determining the crystal structure, phase content, crystallite size, lattice parameters, and other information of the material and the accuracy of the analysis is dependent on the quality of the data and the selected instrumental configuration. the package is compatible with all diffraction scattering techniques that produce powder diffraction data, including the most commonly used laboratory x-ray diffractometer.

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the x-ray diffraction analysis of the samples a, b and c, the three samples obtained from the al-doped synthesized sample has been done and the spectral results are tabulated in table 2. the x-ray diffraction pattern of sample a, b and c, the three samples obtained from the al-doped synthesized sample are shown in figure 1, 2 and 3 respectively. the spectrum pattern obtained from the x-ray powder diffraction has been analysed eliminating the k values and then the peak has been searched using the software x-pert pro ver 3.0. phase identification was performed using the software, search match ver 3.01, from philips and the pdf database file version 2003 from icdd. the x-ray diffraction pattern of the sample a, b and c

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