

Synthesis and AB INTIO Structure Determination Of Mixed Valence a Novel Ternary Oxide $Cd_{0.54} La_{2.5} Pb_{1.25} N_{0.21} O_{0.25}$ and Study the Physical Properties

Yuv Raj Sahu, Rohit K. Dev, Kalpana Mishra and Parashuram Mishra*,
Bioinorganic and Materials Chemistry Research Lab, Tribhuvan University, M.M.A .M. Campus, Biratnagar, Nepal

Abstract

Powder X-ray diffraction (XRD) has been exploited to establish the structural properties of a porous interpenetrated metal framework material prepared by solid-state grinding, recognizing that product phases from mechanochemical synthesis are typically microcrystalline powders. The importance of subjecting the powder XRD data to rigorous scrutiny in such applications is emphasized. The present work deals with the ab initio structure determination of the heavy metal framework having mixed valence in $Cd_{0.54} La_{2.5} Pb_{1.25} N_{0.21} O_{0.25}$ from precession electron diffraction intensities. The metal framework of the compound was solved in this investigation via direct methods from $hk0$ precession electron diffraction intensities recorded with a Philips EM400 at 100 kV. A subsequent (kinematical) least-squares refinement with electron intensities yielded slightly improved co-ordinates for the 11 heavy atoms in the structure. Chemical analysis of several crystallites by EDX is in agreement with the formula $Cd_{0.54} La_{2.5} Pb_{1.25} N_{0.21} O_{0.25}$. Moreover, the structure was independently determined by Rietveld refinement from X-ray powder data obtained from a multi-phasic sample. The compound crystallises in the triclinic crystal system space group $P-1$ with refined lattice parameters of $Cd_{0.54} La_{2.5} Pb_{1.25} N_{0.21} O_{0.25}$ are and refine values $R_p=0.2501, R_{wp}=0.3065$ and $GOF=0.0135$.

Keywords: crystal, structure triclinic, investigation, multiphase, independent

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I. INTRODUCTION

Metal oxides contain mixed valence are very useful for ordering molecules into lamellar structure. These compounds are hybrid inorganic and organic materials and usually consist of alternating inorganic layers. They can be prepared as polycrystalline solids, thin films or membranes depending upon applications. Nowadays, there is great deal of work in metal oxide containing mixed valence due to interesting physical and chemical properties. Recent investigation of structural and conductivity properties of $Cd_{0.775} La_{0.225} O_{1.5}$ oxide conductors (n La, Pr, Nd, Sm, Eu, Gd, Tb, Dy), which belong to the dimorphic rhombohedra $Bi\}Sr\}O$ structural-type family, has proved a close conductivity composition dependence (1). This has been interpreted on the basis of structural data obtained from Rietveld structure refinements based on previous crystal structure investigations (2). The structure is built from cationic slabs parallel to (001) faces of the hexagonal cells. There are nine formula units $La_{0.775} La_{0.225} O_{1.5}$ per triclinic cell, distributed over three slabs. Each slab is constituted from a mixed La^{3+}/n^{3+} layer, sandwiched between two La^{3+} layers, and two oxygen sites are located inside; complementary oxide ions, implied by the formulation stoichiometries, are distributed over one or two sites of the interslab space and exhibit a high mobility, mainly responsible for the conductivity [1]. Depending on the rare-earth nature, a b1 high-temperature form is observed, with a closely hexagonal related structure; its formation from the b2 low-temperature variety occurs during a phase transition that has been attributed to a cationic disordering in the mixed Bi^{3+}/n^{3+} layers. It is accompanied by sudden increases of both lattice parameters, of oxide occupancy in inter slab spaces, and of the conductivity. The pure ion oxide conductor character of the b1 variety has been clearly demonstrated for the alkaline-earth-based solid solutions [2], and has been also verified for lanthanides-based solid solutions. The pure ion oxide conductor character of the b1 variety has been clearly demonstrated for the alkaline-earth-based solid solutions and has been also verified for vanadium-based solid. The thickness of the cationic slabs, which is the largest for the vanadium ternary oxide $Cd_{0.54} La_{2.5} Pb_{1.25} N_{0.21} O_{0.25}$ (the best oxide conductor ever evidenced in this family: at 400°C is $10^{-3} Scm^{-1}$ with $E 0.8 eV$), appears to be an important factor for these attractive conductivity properties. $Cd_{0.54} La_{2.5} Pb_{1.25} N_{0.21} O_{0.25}$ is a term of a wide solid solution domain. The crystal structure of $Bi_{1.25} V_{0.123} Ca_{0.245}$ has been determined from powder diffraction data using a combination of direct methods and the novel approach of applying simulated annealing methods simultaneously to X-ray powder pattern by ab initio method. $Cd_{0.54} La_{2.5}$

$\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ is a polar, non-centrosymmetric, second harmonic generation active material and its crystal structure is one of the more complex to be solved ab initio from powder diffraction data. Many of the important heterogeneous catalysts are transition and non-transition metals dispersed on oxide supports. Although the primary function of the support is to increase the surface area of the metal, there is now a vast literature documenting that supports can cause pronounced changes in the catalytic and chemisorptions properties of the metal [3]. In particular, when group 8-10 metals are dispersed on reducible supports such as oxide of metal ions. The chemisorptions properties and activity of the catalyst depend on the reduction temperature used for pretreatment. Numerous mechanisms have been proposed to account for this suppression of chemisorptions and the associated change in catalytic activity; the most popular is the diffusion of reduced oxide $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ moieties onto the metal, thereby blocking metal sites. However, numerous experimental observations are left unexplained by this model. For example, changes in the magnetic properties of Cd/La, composites with reduction temperature indicate that the Bi does not remain as elemental bismuth, but forms a Cd-La- Pb multiphase phase [3] Unfortunately, investigations into the role of compound formation in producing the observed effect have been hindered by the fact that relatively few ternary oxides containing both an early and a late transition and non-transition metal contained mixed valence are known.. It is interesting to note that superconductivity has been observed in some of the ternary oxide phases having the Bi and V structure type and for $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ In this paper, we report the synthesis and crystal structure of $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ by ab initio method via powder XRD and also study the morphology and electrical property.

Experimental

All chemicals used were analytical grade purchase from Sigma Aldrich U.S.A.. A polycrystalline sample of $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ was synthesized by a standard solid state reaction using a mixture of high purity reagents of $\text{La}_2(\text{CO}_3)_3$ - $\text{Pb}(\text{NO}_3)_2$ and CdCO_3 as the starting materials in the molar ratio of 1 : 1 : 2. The mixture was ground carefully, homogenized thoroughly with methanol (99%) in an agate mortar and then packed into an alumina crucible and calcined at 1000°C in air for 10h with several intermediate grindings [4]. Finally the product was pressed into pellets and sintered at 100 K/h. Powder X-ray diffraction (XRD) data were collected at room temperature in the angular range of $2\theta = 10$ to 90 with scan step width of 0.02° and a fixed containing time of 15 s using Philips powder diffractometer with graphite monochromatic $\text{CuK}\alpha$ radiation. The powder was rotated during the data collection to minimize preferred Orientation effect if any. The program TREOR in CRYSFIRE was used to index the powder pattern which give orthorhombic cell system. SIRPOW92 was used to locate the positional parameters of constituent atoms. The full pattern is fitting and peak decomposition in the space group $Pn-3n$ (222) using check cell program. The structural parameters were refined by the Rietveld method using the JANA program which gave $R_p=0.2501, R_{wp}=0.3065$ and $\text{GOF}=0.0135$. The density is determined by Archimedes principle. The morphology of titled compound was determined using SEM. For the electrical studies, the measurements were preceded by a pretreatment of the sample in order to reduce the mean particle size of the obtained powder. After these treatments, the sample achieved about 85% of the theoretical density with the final diameter of 6 mm and thickness of 23mm. The relative density of the sample before the mechanical grinding was 79 %. Platinum electrodes were connected to the two faces of the pellet via a platinum paste to keep good electric contacts. Impedance spectroscopy measurements were carried out using a Hewlett-Packard 4192a Impedance Analyzer. The impedance spectra were recorded in the 5 Hz-13 MHz frequency range. Electrical conductivity measurements of representative $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ were carried out by complex impedance spectroscopy with a 1174 Solartron frequency response analyzer coupled to a 1266 Solartron electrochemical interface. Pellets of about 14 mm diameter and 1 mm thickness were prepared by cold pressing of a mechanically activated powder mixture with the composition: $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$. To form the phase, the pellets were heated at 700°C during 12 h and slowly cooled to room temperature. This synthesis method was employed to improve the ceramic quality, as it has been shown for other materials [2-6]. The formed phases and crystallinity were studied by X-ray powder diffraction. Platinum electrodes were deposited on the two faces by sputtering, and measurements were carried out in the temperature range 200 - 650°C , at steady temperatures, with pellets under air flow and the frequency range was fixed [5].

Physical Measurements

X-ray powder diffraction data were obtained on a Philips powder diffractometer using a Bragg Brentano geometry, with a back-monochromatized $\text{CuK}\alpha$ radiation having $1.5456(\text{A})$ wavelength.. Diffraction spectra were scanned by steps of 0.023 (2 θ) over the angle range 5 - 50 (2θ), with a counting time of 1.5 s per step. Each sample was rotated 3.14 radian s^{-1} during the data recording, in order to minimize the orientation effects resulting from the material compaction [6]. The pattern peak positions are extracted with the Topas P of Bruker-Diffrac1- 64 software package, where each peak profile is refined using the pseudo-Voigt function. The accurate cell parameters of the different samples were refined from 5 to 50 reflections cubic. The crystal

structure determination of the phase was carried out for the Philips powder diffractometer $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ parent composition. The first attempts were undertaken by recording X-ray diffraction data, but each time, the intensities diffraction pattern was erroneous by the preferential orientation phenomenon. To minimize the preferential orientation effect and indeed be able to separate the oxygen and N atoms of the heavy atoms Bi and V at room temperature. The crystal structure determination has been concluded by using alternately the JANA (16) and EXPO [7] programs. Dilatometric and conductivity investigations were made on samples pelletized at room temperature (diameter 5 mm, thickness approx. 3 mm), then sintered at 900°C for 15 h and quenched to room temperature, in liquid nitrogen. The degree of compactness in all cases ranged between 75 and 92%. Dilatometric studies were performed on a Linseis L75 dilatometer with a heating rate of 0.43°C/min between 20 and 700°C. For conductivity measurements, gold electrodes were vacuum deposited on both #at surfaces of the pellets using the sputtering method. The measurements were obtained by impedance spectrometry in the frequency range 1-106 Hz, using a frequency response analyzer Schlumberger 1170; for a given temperature, each set of values was recorded after a 1h stabilization time. The density was calculated by Archimedes principle.

II. RESULTS AND DISCUSSIONS

The extensive search for novel inorganic materials with open frameworks formed of tetrahedral and octahedral delimiting inter-layer spaces (2D), tunnels (3D) or cages (1D) where cations are housed, represent currently a field of intense activity including several disciplines: solid-state chemistry, physics, mechanics, and mainly ionic conductivity properties and their use as battery materials oxides of metals and alkaline cations are well known for their thermal stability and the simplicity of syntheses, which is important for many practical applications lithium batteries [7]. The compound was prepared from $\text{Bi}_2\text{O}_3\text{-NH}_4\text{VO}_3$ and CaCO_3 as starting material oxides. Calcium oxide powder was dried and decarbonated at 1000 °C over night prior to use. The oxides were weighed in stoichiometric proportions and ground together in an agate mortar. The prepared composition was heated upto 1400 °C for over night in an alumina crucible, no particular condition was used in order to cool down the samples [8]. This procedure was applied at least twice in order to etastabilized compound as verified by X-ray powder diffraction. The final compound is white in colour.

Crystal structure of $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$

The framework structure of $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ was first examined by ab initio structure determination method using the powder XRD data shown in figure 1.. The initial lattice parameters were determined to be parameters and $V=279.7571 \text{ \AA}^3$ by an indexing procedure using the program TREOR15 in EXPO2004.16 The most probable space group was suggested to be cubic crystal system Next, the integrated intensities were extracted by the Le Bail method using the program Jana2006.[9]. A profile function and background function of the Le Bail method used in the present study were pseudo-Voigt function and 20th order Legendre function, respectively. An initial structure model was then obtained by the charge flipping (CF) method17 using the extracted integrated intensities. Although the V site could not be clearly determined by the CF method using the powder XRD data, the framework structure of $\text{Cd}_{0.54}\text{La}_{2.5}\text{Pb}_{1.25}\text{N}_{0.21}\text{O}_{0.25}$ was successfully determined with the help of Rietveld refinement using JANA package shown in figure 2. Rietveld refinement of cubic crystal system having P-1 space group was found with three dimension structure as follows against XRD data shown in figure 3 for structural determination proved difficult, due to a combination of preferred orientation of the plate-like crystallites in flat-plate geometry [10]. In other words, the extracted peak intensities for phase 1 could be reliable. Note that if the relative intensities differed a lot from the simulated patterns of the known phases, additional manual partitioning of those overlapping peaks would be necessary to get reliable intensities for the unknown unknown multi phase 1. After considering the multiplicity and Lorentz-polarization correction, the intensities of phase 1 were used for the structure solution as below. Moreover, the reflection conditions indicated possible space group.P-1. The initial structure model was obtained using a charge flipping algorithm with the program Superflip.8 [11] Random phases were used at the beginning of the charge-flipping interaction, and overlapping peaks were re-partitioned using a histogram match to improve the convergence. The iteration converged with an R factor of 29% and the final electron density shows P1 symmetry with a 5% error. The program of EDMA was then used to automatically assign atomic positions. Four unique heavy atomic positions were found and the heaviest one was assigned as Bi while the others were considered as V and Bi. Due to the existence of heavy atoms, all oxygen positions were ambiguous in the electron density map of this stage[12]. To locate the oxygen atoms, a Monte-Carlo based simulated annealing process with the program TOPAS was applied. For each annealing process, various atomic coordinates were randomly assigned table 2 shown as the initial positions of the oxygen atoms [13]. The annealing process was restarted after finding a few oxygen positions, until all oxygen positions were found to be reasonable with Cd ions. On the whole the crystal structure of cited crystal is octahedral closed face center cubic.

Crystal symmetry

From the analysis of the X-ray powder diffraction data collected on theMPD-PROat1000 °C, we were able to obtain good solutions from two different auto indexing programs Treor [10] and Dicvol implemented in X'Pert High ScorePlus. The solutions determine d by the both programs are very similar: $a=14.1243 \text{ \AA}$ triclinic crystal system, with a figure of merit (FOM) $M_{50}=30$ and $M_{50}=65$, respectively forTreor and Dicvol[13]. From the average oxygen volume in oxides, $V_{\text{oxygen}}=18-22 \text{ \AA}^3$ at room temperature, and the nominal composition $\text{Cd}_{0.54} \text{La}_{2.5} \text{Pb}_{1.25} \text{N}_{0.21} \text{O}_{0.25}$ we can estimate $Z_{\text{max}}=598/18/15=2.39$ and $Z_{\text{min}}=11$. We chose $Z=2$ leading to an oxygen volume slightly larger than the previous values, $V_{\text{oxygen}}=23.3 \text{ \AA}^3$. In order to determine the possible space groups of this compound, an initial procedure was performed to obtain the Bravais class. The observed intensities were extracted using a Le Bail spacegroupP-1.A Patterson function was then calculated with implement in the Fullprof suite. A strong peak was observed at 0.5, 0.5,0 leading to a triclinic Bravais class. In the whole the structure of crystal coordinated by tetrahedral geometry.

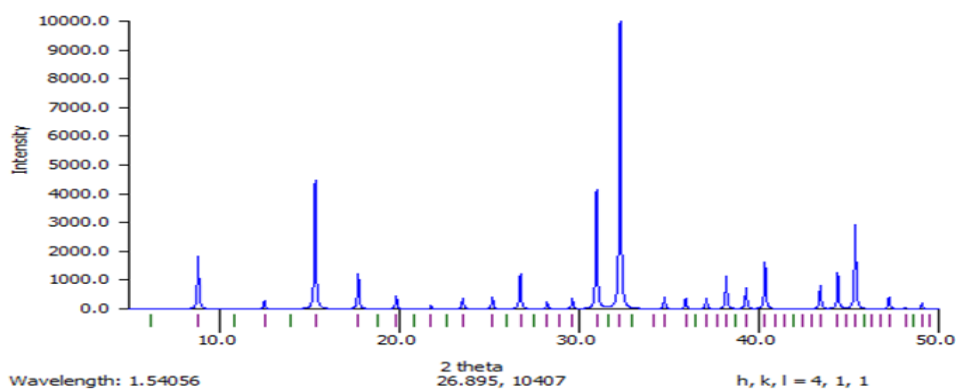


Figure 1 Powder XRD spectra of $\text{Cd}_{0.54} \text{La}_{2.5} \text{Pb}_{1.25} \text{N}_{0.21} \text{O}_{0.25}$

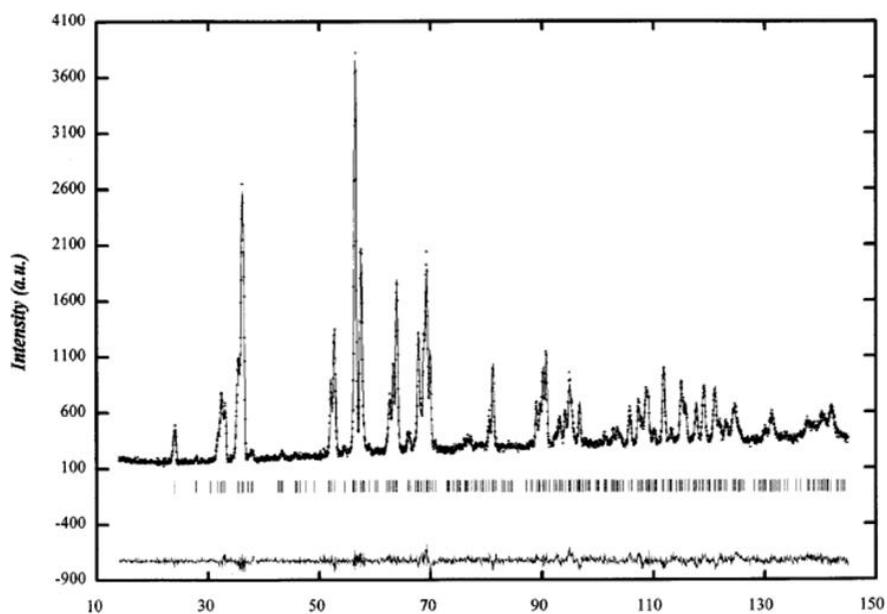


Figure 2 Graphical representation of the result from Rietveld refinement with X-ray powder data. Vertical bars indicate positions of the Bragg reflections for $\text{Cd}_{0.54} \text{La}_{2.5} \text{Pb}_{1.25} \text{N}_{0.21} \text{O}_{0.25}$ dots mark the observed intensities and the solid line gives the calculated intensity curve. The deviations between the observed and the calculated intensities from the refined model are shown by the difference plot in the lower part of the diagram.

Table1.Crystallographic Data

Phase data	
Formula sum	$\text{Cd}_{0.54} \text{La}_{2.5} \text{Pb}_{1.25} \text{N}_{0.21} \text{O}_{0.25}$
Formula weight	673.9 g/mol
Crystal system	triclinic
Space-group	P -1 (2)
Cell parameters	$a=3.1346 \text{ \AA}$ $b=5.3402 \text{ \AA}$ $c=6.0869 \text{ \AA}$ $\alpha=79.9066^\circ$ $\beta=76.8208^\circ$ $\gamma=85.1414^\circ$

Cell ratio	a/b=0.5870 b/c=0.8773 c/a=1.9418
Cell volume	97.57 Å ³
Z	2
Calc. density	22.9378 g/cm ³
Meas.density	22.91671 g/cm ³
Pearson code	aP17
Formula type	N2O2P5Q8
Wyckoff sequence	i8d

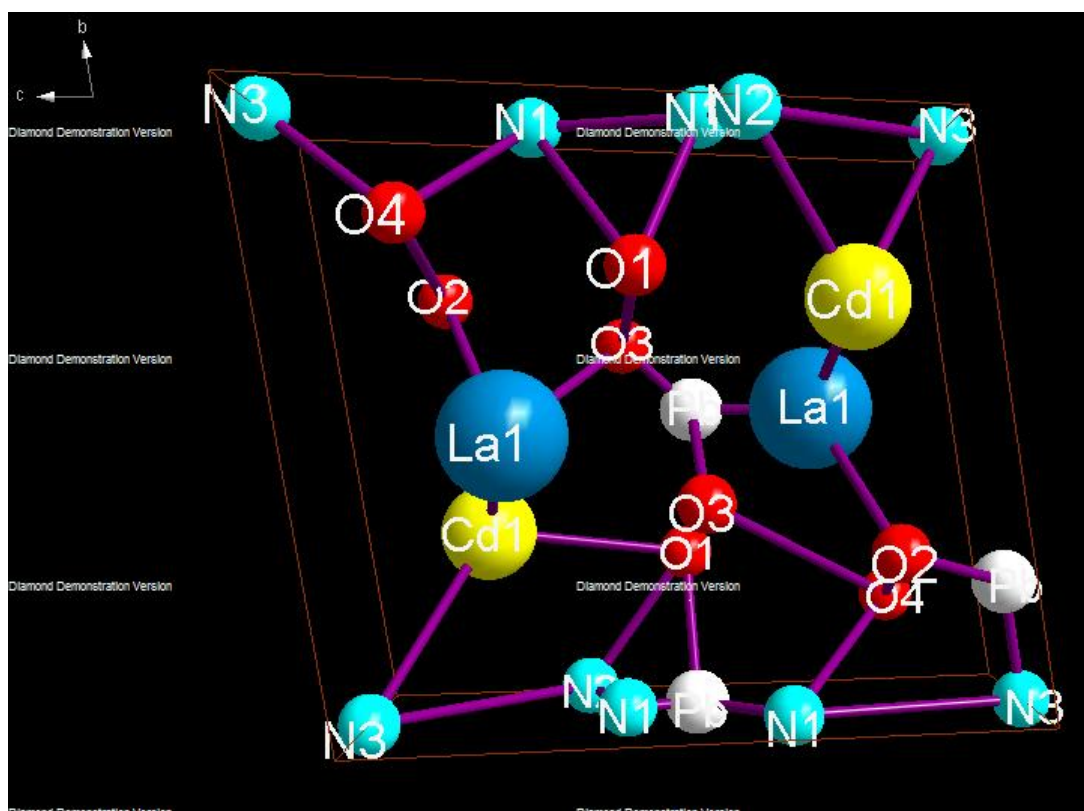


Figure 3. Crystal structure of Cd_{0.54}La_{2.5}N_{0.21}O_{0.25}Pb_{1.25}

Table 2. Fraction Atomic parameters

Atom	Ox.	Wyck.	Site	S.O.F.	x/a	y/b	z/c	U [Å ²]
La1	+3	2i	1		0.39874	0.52349	0.26485	0.0380
Cd1	+2	2i	1		0.89532	0.69952	0.19451	0.0380
O1	-2	2i	1		0.08875	0.25861	0.49731	0.0380
O2	-2	2i	1		-0.09756	0.28617	0.18790	0.0380
O3	-2	2i	1		0.23189	0.63376	0.54389	0.0380
O4	-2	2i	1		0.22731	0.17865	0.17981	0.0380
N1	-2	2i	1		0.63272	0.00103	0.37387	0.0380
N2	-2	2i	1		0.93688	0.98752	0.31186	0.0380
N3	-2	1d	-1		1/2	1.00000	0	0.0380
Pb	-2	2i	1		0.50000	0.21345	0.00000	
Pb	-2	2i	1		0.21450	0.52143	0.45120	
Pb	-2	2i	1		0.32160	0.00000	0.51650	

Table 3. Selected Bond angles

Atom1	Atom2	Atom3	Angle(^o)
La1	Pb	Cd1	112.455
	Pb	O3	22.976
	Pb	Cd1	84.453
	Pb	O3	71.622
	Pb	Pb	133.323
	Pb	O2	120.684
Cd1	Cd1	O3	91.068
	Cd1	Cd1	116.902
	Cd1	O3	71.991
	Cd1	Pb	60.172
	Cd1	O2	71.614
Pb1	Cd1	O1	141.206
	Pb2	O4	134.271
	Cd1	O3	160.593
	Cd1	O1	105.024
	Cd1	O2	154.297
	Cd1	Pb	137.504
O1	Cd1	Pb	64.653
	Cd1	Pb	107.056
	O3	Cd1	86.547
	O3	O3	70.513
	O3	Pb	115.954
	O3	La4	123.396
	O3	O1	69.405
	O3	O4	127.043
	N1	O3	69.526
	N3	O1	31.253
	O3	O2	114.379
	O3	Pb	46.707
	O3	Pb	52.088
	N4	Pb	155.698
	Cd1	O3	155.951
	Cd1	Pb	64.892
	Pb5	Cd1	O2
Cd1		O1	95.734
Cd1		O4	92.394
O3		Pb	93.452
O3		Pb	22.234
O3		Pb	99.311
Pb		O2	101.301
Pb		O1	158.506
Pb		O4	111.068
Pb		O3	127.819
Pb		O1	98.486
Cd1	Pb	Pb	121.289
	La1	N2	122.772
	La1	La1	116.902
	La1	Pb	62.936
	La1	O1	97.696

The heavy-atom positions M have been refined, where as the positions of the oxygen atoms have been transformed from the structural model suggested by Lundberg and Sundberg [14]. Each M site was assumed in the refinement as occupied by 0.49 Cd and 0.51La. The calcium sites refined to an occupancy of 0.648(10) for Ca-1 in the 6-sided tunnels and 0.392(8) for Ca-2 inside the 7-sided tunnels. Lattice parameters for the a and b axes, determined from a calibrated X-ray diffraction pattern recorded along the [0 0 1] direction, are a =14.1243, b =14.1243, c=14.1243 Reflection intensities I_{h k l} were estimated using the crysfire program . Merging the intensities according to sym- symmetry yielded a data set of 556 crystallographically non-equivalent hk0

reflections after removal of 55 symmetry-forbidden reflections with non-zero intensity. The internal R-factor of symmetry-related reflections in the merged data set was 15.8% with a resolution. According to our analysis space group $Pb3n$ was tried for solving the structure by direct methods with the program SIR97 [15]. The program was modified for this purpose with electron scattering factors provided by Jiang and Li . Similar to previous studies] suspected dynamical diffraction was taken into account by using a phenomenological compensation based on The heavy Bi and V atoms, which are shown in the figure by the large grey shaded spheres, form with the oxygen atoms (small red spheres) the structural fragments of MO6 octahedra respectively. The calcium atoms are located inside the 7- (Cd-1) and 6-sided tunnels (Cd -2). The positions of the oxygen atoms have been taken from the proposed structural model of Lundberg and Sundberg and transformed to cubical space group $Pn-3n$. The approximation $|hkl|/|Vhkl|$ that has been derived as the limit for thick crystals under two beam conditions, where $Vhkl$ (in volts) designates the structure factor amplitude for electrons. All attempts to use $|h k l|/|Vh k l|^2$ instead led to wrong results with direct methods and least-squares refinement. Moreover, we found that performed geometrical corrections for parallel illumination, as suggested by other authors [16], spoiled the data with the result that it was neither possible to solve the structure nor to refine a given model to good geometry. Following our results obtained for the Cd_{0.54} La_{2.5} N_{0.21} O_{0.25} Pb_{1.25} nanocrystalline powder, the data were used throughout this study without correction. The most probable solution found with these data by SIR97 had an R-factor of 40%. All 40 heavy atoms in the unit cell are clearly visible in the potential map that was calculated by the program using the 95 largest $hk0$ E-values [16]. Coincidence of several additional weak peaks in the map with suspected oxygen or cadmium atoms proved accidental and was not considered as reliable. A (kinematical) least-squares structure refinement on $|Vh k l|$ was performed for the detected 11 heavy-atom positions in the asymmetric unit (program SHELXL97-2. For the refinement, it was finally assumed that the heavy-atom positions are fully occupied by tungsten because only this yielded positive defined temperature factors for all atoms. This was not the case if mixed Nb/W sites were assumed and it is an indicator for the presence of dynamical effects. The overall R-factor of the refined framework structure was 35.2% (35] parameters refined, unit weights). The refined atomic co-ordinates together with the displacement factors U_{eq} are listed in Table 3. The average positional shift of the atoms during the refinement was 0.07 \AA (maximum shift 0.11 \AA for atom M5). The potential map after least-squares refinement from X-ray diffraction is shown in figure 1. An attempt to detect the oxygen positions from a V_{obs}/V_{calc} difference Fourier map was unsuccessful due to the relatively high background level (lowest peak to background ratio about 3). Although several diffuse peaks appeared at the expected positions for calcium inside the 6-sided and 7-sided tunnels, the quality of the data did not allow reliable location of these atoms. Thus, the full range of available intensities up to resolution of 0.73 \AA^{-1} was tested. Refinement of the heavy-metal framework with 651 unique reflections and mixed occupancies (Bi/V = 1.0) yielded an R-factor of 41.6 %. However, despite several additional peaks appeared in the $V_{obs}-V_{calc}$ differential Fourier map, it was again not possible without knowing the correct positions in advance to assign these peaks to the missing atoms. Since the quality of the received framework model could not significantly be improved by the data with higher resolution, we present here only the result obtained with intensities up to 1 \AA^{-1} . Nevertheless, in order to check the effect of geometrical correction on the quality of the data, we calculated the required correction factors f_c which match theoretical structure factors $V_{h k l, calc}$ with uncorrected structure factors $V_{h k l, exp, raw}$ and geometrical corrected structure factors $V_{h k l, exp}$ precess corrected, respectively: $f_c = V_{h k l, calc} / V_{h k l, exp}$; $f_c = V_{h k l, calc} / V_{h k l, exp}$; raw.

Crystal structure description

The crystal structure of this compound can be described by different cationic environments. Each mixed site V/Bi(1) is surrounded by eight oxygen atoms (six O(1) atoms and two O(2) atoms) at normal distances ranging from 2.47 to 2.57 Å with a mean distance O/Ca 2.51 Å to form a slightly distorted cubical polyhedron (Fig. 4). This high symmetry of the oxygen environment leads to a very weak stereochemical activity and the nonhybridization of the La (1) lone pair (Lp) of the atom, which is likely very close to the bismuth nucleus. The second atom La(2) is bonded by seven oxygen atoms to form a very distorted polyhedron, with two very short distances (2.05 and 2.09 Å for respectively the O(3) and O(1) atoms) and normal distances varying from 2.31 to 2.49 Å. This distorted environment is often observed when a Bi atom possesses a lone pair and the refinement of the Bi(2) electronic lone pair Lp2 places it 1.22 Å from the nucleus, to occupy the interlayer between adjacent Bi(2) Ca layers . Each V-Bi(1) atom shares O(1) atoms with the six neighboring cations V-Bi(1) to form a mixed Ca-Bi-O layer parallel to the plan (b, c), associated with angles shown in table 4. In the same manner, each Bi(2) atom shares also oxygen atoms with the six closest bismuth atoms, to form a layer parallel to the precedent (Fig. 4b). Thus, the crystal structure can be described by a piling of layers ABC, ABC, etc., in the a-axis direction, where layers A and C are formed by Bi(2) atoms and their oxygen environment, and layer B is constituted by mixed sites La}Bi(1) and their oxygen atoms (Fig. 5). The cohesion between layers A and B is ensured by the O(1) and O(2) atoms with long distances to the Cd-La-Pb(1) atoms (Fig. 4) and all bond angles are shown in table 4.

Morphological study by SEM (scanning electron microscopy)

Scanning electron microscopy (SEM) is giving morphological examination with direct visualization. The techniques based on electron microscopy offer several advantages in morphological and sizing analysis; however, they provide limited information about the size distribution. For SEM characterization, nanoparticles solution should be first converted into a dry powder, which is then mounted on a sample holder followed by coating with a conductive metal, such as gold, using a sputter coater [17]. The sample is then scanned with a focused fine beam of electrons. The surface characteristics of the sample are obtained from the secondary electrons emitted [18] from the sample surface. The morphology of the oxide nanoparticles is shown in Fig.2. From the image, it is clear that the particles were highly agglomerated in nature. The SEM pictures clearly show randomly distributed grains with smaller size. From the SEM analyses, one can conclude the formation of nanoparticles spherical structure. Here it is grown in very high-density and possessing almost uniform spherical shapes. The image reveals that the average size of the particles is 40.32 nm.

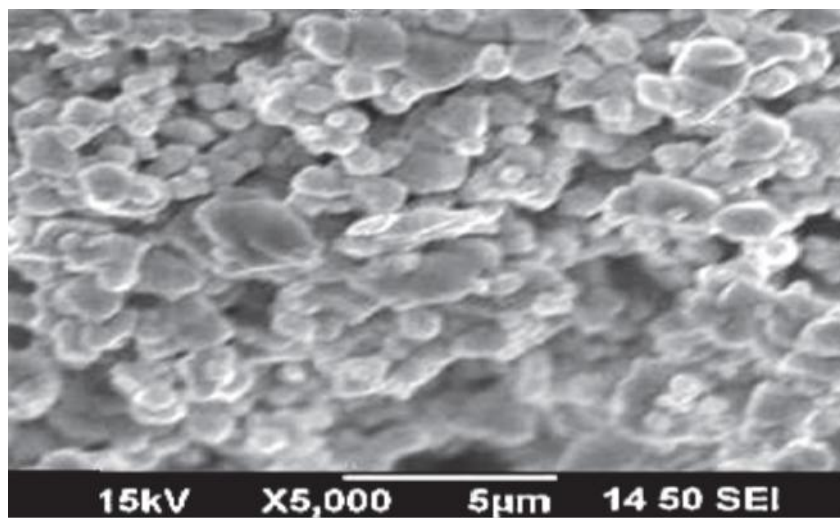


Figure 5 SEM spectrograph of $Cd_{0.54} La_{2.5} Pb_{1.25} N_{0.21} O_{0.25}$

The sample is then scanned with a focused fine beam of electrons. The surface characteristics of the sample are obtained from the secondary electrons emitted [12] from the sample surface. The morphology of the oxide nanoparticles is shown in Fig.2. From the image, it is clear that the particles were highly agglomerated in nature. The SEM pictures clearly show randomly distributed grains with smaller size. From the SEM analyses, one can conclude the formation of nanoparticles spherical structure. Here it is grown in very high-density and possessing almost uniform spherical shapes. The image reveals that the average size of the particles is 40.32 nm.

Electrical properties

The electrical properties of the title compounds were determined by using complex impedance spectroscopy. The electrical measurements were carried out in air in the 240–360°C temperature range after stabilization at each step of 30 °C and in the frequency range 1Hz-13MHz. The applied voltage was 0.5 V, which allows eliminating aberrant points at low frequencies. Z-view computer program [19] was used to determine the electrical parameters by using a conventional electrical circuit as follow: R//CPE-R//CPE, where CPE is a constant phase element:

$$Z_{CPE} = \frac{1}{A(j\omega)^p} \dots\dots\dots 1$$

An additional inductance L was added to account for instrumental contributions especially at high temperature. The true capacitance was calculated from the pseudo-capacitance according to the following relationships

$$\omega_0 = (RA)^{-1/p} = (RC)^{-1} \dots\dots\dots 2$$

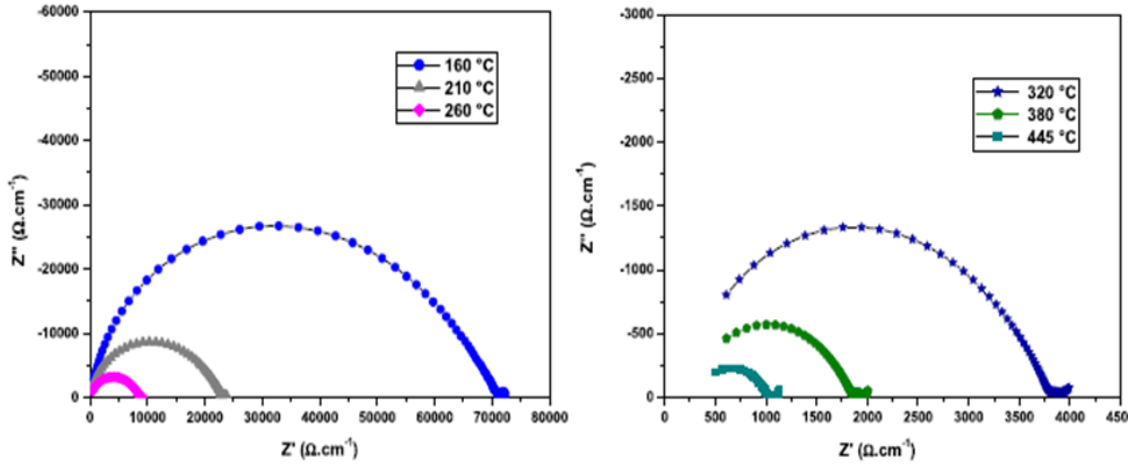


Figure 6. Normalized impedance spectra recorded on Bi_{1.25} V_{0.123} Ca_{0.245} N_{1.24} O₈ at 160 °C–445 °C.

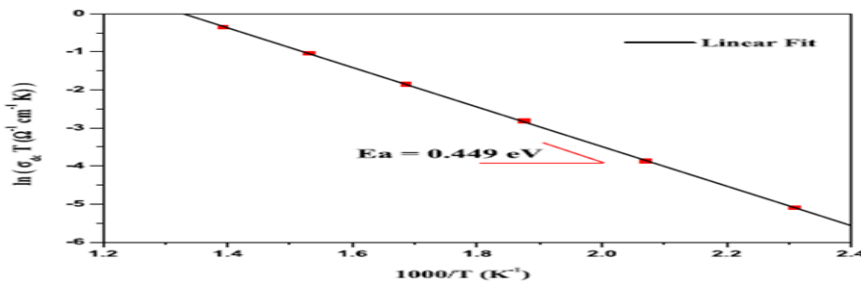


Figure 7 Arrhenius plots of conductivity of Bi_{1.25} V_{0.123} Ca_{0.245} N_{1.24} O₈.

Table 4. Electrical values of the equivalent circuit parameters calculated for Bi_{1.25} V_{0.123} Ca_{0.245} N_{1.24} O₈ sample at different temperatures.

T(°C)	T(K)	C ₁ (×10 ⁻¹¹ F) ± 0.02	P ₁ ± 0.02	ρ(kΩ.cm) ± 0.01	σ(×10 ⁻⁴ S.cm ⁻¹) ± 0.01
160	433	1.49	0.9	78.832	0.14
210	483	1.51	0.9	23.263	0.43
260	533	1.64	0.9	8.958	1.11
320	593	1.59	0.9	3.788	2.64
380	653	1.58	0.8	1.852	5.40
445	718	1.70	0.8	1.022	9.78

Geometric value of pellet g = 0.812 cm⁻¹.

normalized impedance spectra recorded on Cd_{0.54} La_{2.5} N_{0.21} O_{0.25} Pb_{1.25} at 150 °C–445 °C are shown in figure 6. The electrical parameters values achieved from the equivalent circuit in the temperature range 160–445 °C are summarized in table 3. Where, the resistivity $\rho = R/k$ is extracted from the refinement of each contribution which the geometric factor of the cylindrical pellet $g(\text{cm}^{-1}) = e/S$ (e =thickness; S =surface). The conductivity increases from $0.14 \times 10^{-4} \text{ Scm}^{-1}$ at 160 °C to $9.78 \times 10^{-4} \text{ Scm}^{-1}$ at 445 °C (table 3). Although, the absolute conductivity value of the Al/Li-substituted title material at 320 °C ($\sigma = 2.64 \times 10^{-4} \text{ Scm}^{-1}$). The Arrhenius plot of the electrical conductivity, $\log(\sigma T (\text{S.Kcm}^{-1}))$ as a function of $1000/T (\text{K}^{-1})$, in the temperature interval 160–445 °C is illustrated in figure 7. As a single linear plot and following the Arrhenius law, the activation energy of the Bi_{1.25} V_{0.123} Ca_{0.245} N_{1.24} O₈ compound determined by linear fit is 0.549 eV and shows pseudo-order of kinetic thermal reaction. Its conductivity increases with increasing temperature. This material presents interesting electrical performance and can be used as super-conductor materials [19].

III. CONCLUSIONS

The title compound, Cd_{0.54} La_{2.5} Pb_{1.25} N_{0.21} O_{0.25} has been synthesized as polycrystalline powder by solid-state method. To the present authors' knowledge, our work is the first to uncover the metal framework structure of a large unit cell heavy-metal oxide contained mixed valence with tunnel structure directly from single-projection X-ray diffraction data. To reach this goal, large-angle precession X-ray diffraction intensities have been collected on imaging plates at 100 kV. Processing of the data was performed following the quasi-kinematical approach as outlined elsewhere. Suggested data correction schemes for the geometry of off-axis hollow-cone illumination have been checked, but were not applied because this impaired the quality of the data. Thus, the framework structure of the title compound was solved from uncorrected data with 1 Å resolution using the direct methods program SIR97. All atoms in the asymmetric unit of the Cd_{0.54} La_{2.5} N_{0.21} O_{0.25} Pb_{1.25} framework were readily found by the program and correctly assigned with their 3D coordinates. A (kinematical) least-squares structure refinement of the heavy-atom positions was carried out with the program JANA using the same hkl data set. Comparison of the refined co-ordinates for the heavy atoms resulting from Rietveld refinement on X-ray powder data (present study) showed an average agreement for the metal framework structure within 0.09 Å. SEM analysis of several crystallites examined indicate an average chemical composition of this phase according to Cd_{0.54} La_{2.5} N_{0.21} O_{0.25} Pb_{1.25}. Despite the quality of the available precession X-ray diffraction data proved sufficient for accurate determination of the heavy-atom positions, the data quality did not allow to unambiguously locate the vanadium atoms inside the tunnels nor to reveal the oxygen atoms. Theoretical calculations carried out for the M5O14 framework structure show that simple formulas for the detectability of light atoms in the presence of heavy atoms can be misleading, since the background noise which results from truncation errors in the Fourier synthesis is not taken into account. Thus, for calculated data with 1 Å resolution it was found that the heavy atoms appear effectively 7.8 times stronger in the projected map than the oxygen. This explains why it would be very difficult with real data to detect the oxygen and nitrogen positions, in particular if only data from a single zone axis are considered as in the present case. The fact that it was neither possible to detect the oxygen nor the calcium atoms in the structure is a strong indicator that the data from precession X-ray diffraction still contain considerable amounts of dynamical/secondary scattering. This is also displayed by the relatively large R-factors after LS-refinement (40.2%). From our point of view, the fact that it was possible to determine the heavy-atom positions with high precision even from 100 kV data is yet a great success, because solving these types of heavy-metal framework structures directly from X-ray diffraction data was attempted by several groups for many years, but obviously, without success. Therefore, the present result is a striking example where precession X-ray diffraction reduced dynamical diffraction below the critical limit at which the determination of the structure or, at least, a fundamental part of it is enabled. However, further experiments which address to optimized conditions for detecting the oxygen atoms in heavy-atom structures seem to be necessary. So collection of 3D intensity data are likely more promising to reach this goal, since such extended data sets proved sufficient to detect even the most light atom, oxygen and nitrogen atoms. So from the above discussion we concluded that the structure of cited compound by powder pattern and the particle show nanomaterial can be used in super conductor.

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