



**HAL**  
open science

## Dielectric properties of colossal permittivity materials: An update

Chafé Cheballah, Zarel Valdez Nava, Lionel Laudebat, Thierry Lebey, Pierre Bidan, Sombel Diaham, Sophie Guillemet-Fritsch

► **To cite this version:**

Chafé Cheballah, Zarel Valdez Nava, Lionel Laudebat, Thierry Lebey, Pierre Bidan, et al.. Dielectric properties of colossal permittivity materials: An update. *Journal of Energy and Power Engineering*, 2013, vol. 7 (n° 4), pp. 726-732. 10.1109/CEIDP.2011.6232754 . hal-01169490

**HAL Id: hal-01169490**

**<https://hal.science/hal-01169490>**

Submitted on 29 Jun 2015

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



## Open Archive TOULOUSE Archive Ouverte (OATAO)

OATAO is an open access repository that collects the work of Toulouse researchers and makes it freely available over the web where possible.

This is an author-deposited version published in : <http://oatao.univ-toulouse.fr/>  
Eprints ID : 14110

**To cite this version** : Cheballah, Chafe and Valdez-Nava, Zarel and Laudebat, Lionel and Lebey, Thierry and Bidan, Pierre and Diahm, Sombel and Guillemet-Fritsch, Sophie Dielectric properties of colossal permittivity materials: An update. (2013) Journal of Energy and Power Engineering, vol. 7 (n° 4). pp. 726-732. ISSN 1934-8975

Any correspondence concerning this service should be sent to the repository administrator: [staff-oatao@listes-diff.inp-toulouse.fr](mailto:staff-oatao@listes-diff.inp-toulouse.fr)

# Dielectric Properties of Colossal Permittivity Materials: An Update

Chafe Cheballah<sup>1,2</sup>, Zarel Valdez-Nava<sup>1,2</sup>, Lionel Laudebat<sup>3</sup>, Thierry Lebey<sup>1,2</sup>, Pierre Bidan<sup>1,2</sup>, Sombel Diahm<sup>1,2</sup> and Sophie Guillemet-Fritsch<sup>4</sup>

1. LAPLACE (Laboratoire Plasma et Conversion d'Energie), Université de Toulouse, Toulouse 9 F-31062, France

2. CNRS (Centre National de la Recherche Scientifique), LAPLACE, Toulouse F-31062, France

3. Centre Universitaire Jean-François Champollion, Albi 81012, France

4. Institut Carnot CIRIMAT (Centre Inter-universitaire de Recherche et d'Ingénierie des Matériaux), Université Paul Sabatier, Toulouse 9 31062, France

**Abstract:** During the last 10 years, a lot of interests have been devoted to the so-called CDC (colossal dielectric constant) materials. The first materials exhibiting this behavior were the perovskite-based ceramics on the CCTO ( $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$ ) composition. Relative dielectric permittivity can attain values up to (or even larger than)  $10^5$ . Nevertheless, their dielectric losses are too high, the lower values ranging 10%, in a narrow frequency range, thus limiting their applications. The underlying physical mechanisms at the origin of the CDC are still under study. The analysis of broadband impedance spectroscopy measurements leads most of the authors to propose an interfacial polarization mechanism (at the electrodes or at internal barriers), there is a limited number of complementary electrical characterization techniques, which, up to now, comfort the proposed interfacial polarization mechanisms. In the present work, I-V and time-domain polarization are used to characterize these materials. One of the main results is the observation of a non-symmetrical response of these materials related to the direction of the polarization. These results are observed for both macroscopic level on bulk polycrystalline material and within individual grains of the same samples. These results do not fit current accepted models for polarization for CDC materials.

**Key words:** Colossal permittivity,  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$ , CCTO dielectric properties.

## 1. Introduction

Materials presenting the so-called colossal dielectric permittivity continue to attract a lot of attention since they were observed in CCTO ( $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$ ) ceramics [1]. These show a relative dielectric permittivity ( $\epsilon_r$ ) of up to  $10^5$  (at room temperature) that remains constant over a large temperature range with no apparent ferroelectric transition or phase transformations below 732 K [2]. These interesting properties that are stable with temperature make these materials suitable

candidates for miniaturized capacitive elements in microelectronics. Despite the intensive works carried out to understand the physical origins of the high dielectric permittivity in polycrystalline ceramics [3], or in single crystals [4], or in thick and thin film layers [5], the exact role of both composition and microstructure on the electric properties of CCTO are still under study.

The earliest works intended to relate an intrinsic polarization to the crystalline structure. After inconclusive results [6], an extrinsic origin was proposed, assuming that interfaces were the key parameter on the polarization mechanism of CCTO.

First, these interfaces can be associated to the contact of the material to the electrodes [7], where the

---

**Corresponding author:** Zarel Valdez-Nava, Ph.D., CNRS researcher, research fields: electroceramics, high-temperature ceramic dielectrics and colossal permittivity phenomena. E-mail: valdez@laplace.univ-tlse.fr.

differences in the metal work functions modify the dielectric response of the electrode-ceramic-electrode assembly. Supporting this hypothesis, Deng et al. [8] observed significant modification on the  $I$ - $V$  (current-voltage) characteristics related to the surface treatments on the samples. On the other hand, the characterization of polycrystalline CDC (colossal dielectric constant) materials by means of BIS (broadband impedance spectroscopy) identifies at least two distinct responses by equivalent circuit modeling [9]. It was suggested that one response corresponded to semiconducting grains, and the other to insulating grain boundaries. This analysis opened the way to another intrinsic barrier-induced polarization scheme, where the internal barriers formed by the grain boundaries acted as in IBLC (internal barrier layer capacitors). BIS remains the most used characterization technique for CDC materials [4, 9-13].

Only a limited number of studies using other characterization methods than BIS analysis in polycrystalline CDC materials have been reported. Local current-voltage ( $I$ - $V$ ) measurements using microelectrodes and local probe on polished CCTO confirmed the IBLC hypothesis, when Chung et al. [14] measured a barrier height at the grain boundary with a strong non-linear response. Scanning impedance microscopy and C-AFM (conductive atomic force microscopy), showed an insulating secondary-phase inclusion within a single CCTO grain [15]. Recently a controversy appeared [16], when Fu et al. [17] found that grain boundaries were also semiconducting and that grains were a mixture of semiconducting and insulating regions.

Further complementary characterization performed on colossal permittivity materials allowed to identify non-symmetrical response related to the applied voltage that is not taken into account by current accepted models [18]. In the present work, CCTO ceramics were analyzed by additional electrical characterization techniques, such as time-domain polarization at the bulk samples and  $I$ - $V$  characterization, both for the bulk and

by locally probing the characteristics at the grain and grain boundary scale. Such studies were undertaken in order to add further elements to the discussion of the mechanisms involved in CDC materials polarization.

## 2. Experimental Procedure

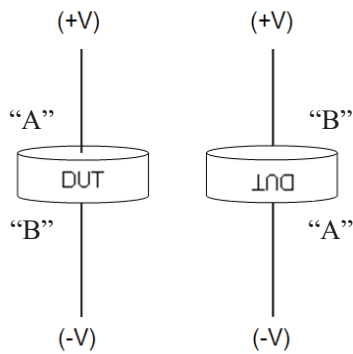
The powders were prepared using a soft chemistry method described previously [19]. They were then pressed into pellets (5 mm diameter and 1.2 mm thickness) under 250 MPa and sintered in air at 1,050 °C for 24 h. The surfaces of the CCTO sample were polished (final thickness, 1.1 mm) and sputtered gold was applied as electrode.

Impedance spectroscopy measurements confirmed that the apparent permittivity of the samples presents the previously described characteristics of CDC materials [9, 20, 21].

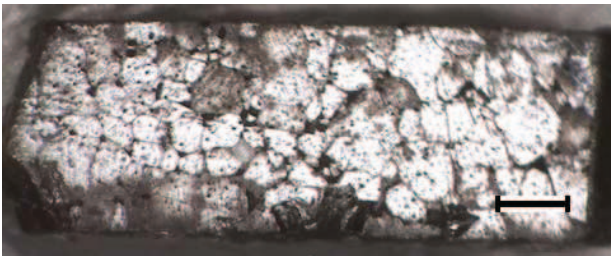
In order to determine the influence of the electric field orientation on the electric response of the sample, each electrode was clearly identified “A” and “B”. A Keithley 2410 sourcemeter is used for  $I$ - $V$  measurements, and in the measurement set-up, when the positive terminal of the sourcemeter is connected to the electrode on the sample surface “A”, the position is said to be “A (+)” and “B (+)” in the opposite case (Fig. 1).

To perform local  $I$ - $V$  characterizations at the grain and grain boundary scale, the CCTO pellet is sawed with a diamond wire cutting machine (ESCIL, W3241 model). The section was polished and thermally etched at 950 °C for 20 min to reveal the grain boundaries for optical observation. Scanning Electron Microscope (SEM-FEG, Jeol JSM 6700-F) coupled with an Energy Dispersive X-ray detector (EDX, Princeton Gamma Tech) and optical microscopy observations show large grain size distribution (up to 150  $\mu$ m) with some pores at the surface (Fig. 2).

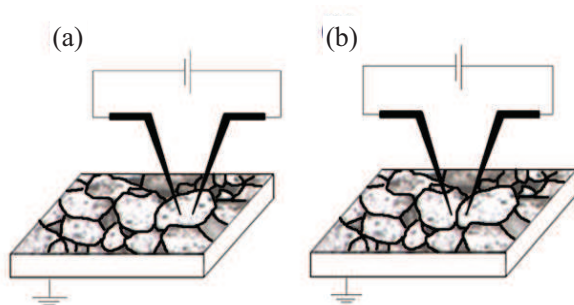
A thermo-regulated, microprobe station (Signatone S-1160 model) with tungsten probes (10 $\mu$ m in radius) is used to ensure local contact on the surface of the grains at 25 °C (Fig. 3). The two tungsten probes are connected to the Keithley 2410 sourcemeter.



**Fig. 1** Definition of the two possible configurations for the sample.



**Fig. 2** SEM micrograph of CCTO ceramics sintered at 1,050 °C for 24 h (bar 500 μm).

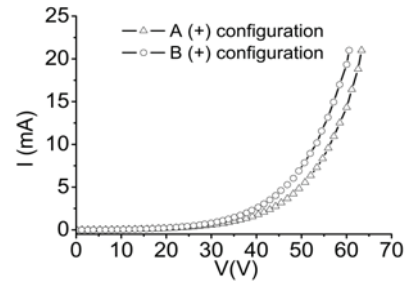


**Fig. 3** Microprobes configuration for  $I(V)$  measurements (a) within the grain and (b) across grain boundaries.

For time domain current measurements ( $I-t$ ) a Keithley 6517A apparatus is used. Voltage steps are applied for polarization ranging from 0.1 V to 20 V. During the short-circuit of the sample, depolarization current is recorded.

### 3. Results and Discussion

Fig. 4 shows the  $I-V$  measurements at room temperature. A non-linear characteristic is observed [14, 22]. However, when superposing the  $I-V$  responses of the “A (+)” and “B (+)” relative to the positive terminal of the sourcemeter, a non-symmetrical response of the sample is observed.



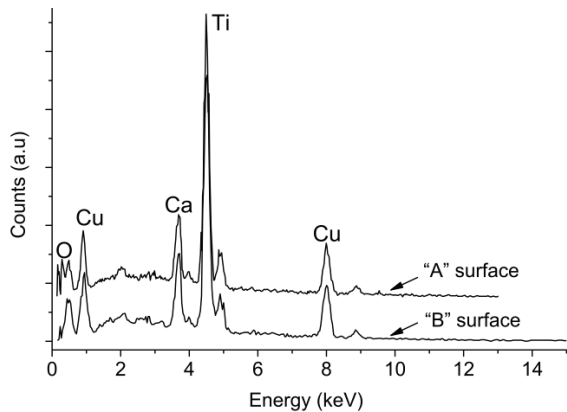
**Fig. 4** Bulk  $I-V$  measurements for the CCTO ceramics in “A (+)” and “B (+)” configurations at room temperature.

These data are reproducible, either by changing the polarity of the applied voltage, when changing the wiring configuration, or when sample is turned upside down in the measured device cell. No memory effect can subsist since the sample is short-circuited for 5 min between every measurement (the current reaches a value of 0.2 pA).

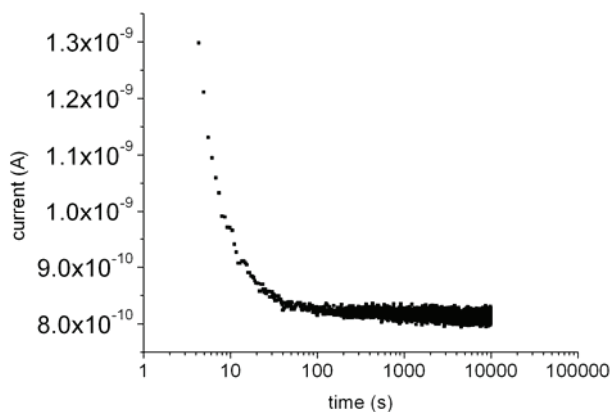
The non linear  $I-V$  response has already been treated in other work and is assumed to be related to a Schottky contact allowing a potential barrier at the interface between grain and grain boundary [12, 23] or between the metal and the material [13, 24]. If the material is considered to be homogenous, only the interfaces between the CCTO ceramic and the metal electrode can explain the non-symmetrical phenomena. To check this assumption, the compositions determined from EDX analysis from each of the two sides (“A” and “B”) to the middle of the pellet were compared. Figs. 5a and 5b show that there is no significant difference in elemental composition of the pellet. So, if the chemical composition is homogeneous, the electrode contact can be one possible source for the difference of the electrical responses between the “A” and “B” configurations.

Deng et al. [5] have already observed this difference when applying platinum electrodes on thin-film CCTO with different PLD (pulsed laser deposition) conditions.

Fig. 6 shows  $I-t$  measurements for very low applied voltage (0.1 V). A quasi steady-state current is reached after  $10^2$  s. In conventional dielectric materials, when a step voltage is applied, the current evolution as function of time is composed of two phases:



**Fig. 5 Elemental composition (EDX) from surfaces “A” and “B” at the center of the pellet.**



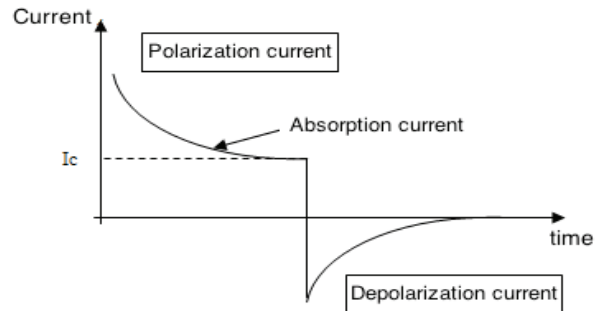
**Fig. 6 Polarization current for the CCTO ceramic with an applied voltage step of 0.1 V.**

polarization and depolarization currents, as represented in Fig. 7. The polarization current is also composed of two parts, the absorption current mainly due to the polarization of dipoles in the dielectric, which dominates at short time, whereas the second part of the curve corresponds to the conduction current ( $I_c$ ). The absorption current of the CCTO sample (without  $I_c$  contribution) and the absolute values of the depolarization current are super-imposable.

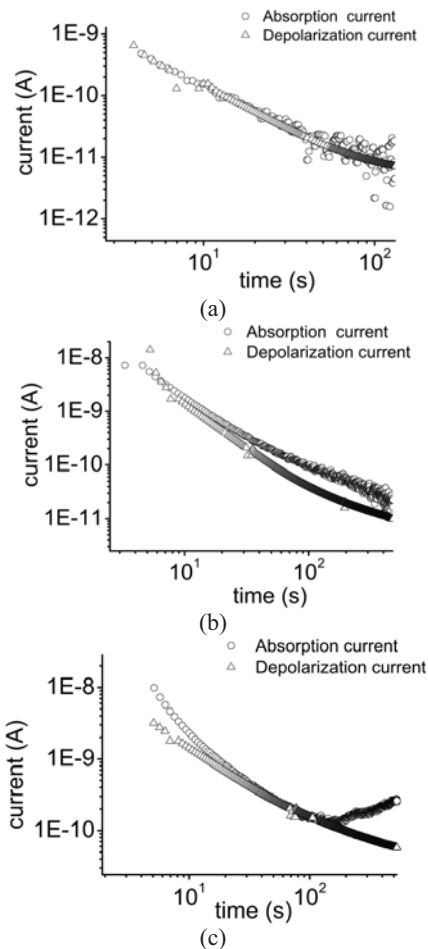
This behavior means that the stored charges in the material during polarization are restored during the depolarization.

In CCTO bulk sample, when absorption and depolarization currents are compared at low voltage, a dielectrical-like behavior as observed (Fig. 8a). This behavior exists up to 2 V (Figs. 8b and 8c).

For higher applied voltages ( $V > 2$  V), the currents are no longer super-imposable (Fig. 9).



**Fig. 7 Classical evolution of time-current response (Polarization and Depolarization) after an applied step voltage to a dielectric material.**



**Fig. 8 I-t polarization response of bulk CCTO at ambient T and (a)  $V = 0.1$  V; (b)  $V = 1$  V and (c)  $V = 2$  V.**

The very high values of the conduction current are shown in Fig. 10 (0.3 mA for 5 V), they are related to the strong non-linear  $I$ - $V$  behavior of CCTO.

When probing the  $I$ - $V$  response at a local level, care should be taken when interpreting surface measurement results, as the exact current path and the

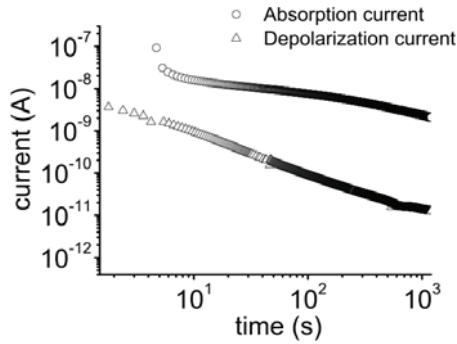


Fig. 9  $I-t$  response comparison between absorption and depolarization currents of bulk CCTO at room temperature and  $V = 5$  V.

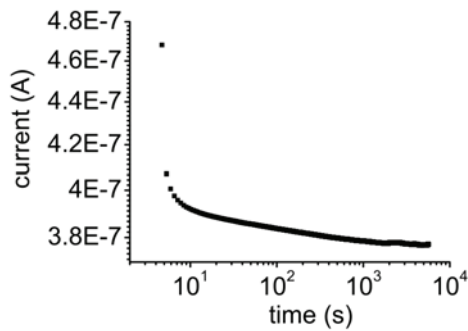


Fig. 10  $I-t$  polarization response of bulk CCTO at ambient  $T$  and  $V = 5$  V.

field distribution are unknown. The  $I-V$  responses of the grains and of the grain boundaries (Fig. 11) are both strongly non-linear. Furthermore, when reversing the voltage direction, thus, the current direction also, a non-symmetrical response is observed. Although different current values were observed between grain and across grain boundaries, due to the complexity of the measurements, only attention is given to their non-symmetrical behavior relative to the current direction.

When performing  $I-V$  probes measurement through gold electrodes contact, Chung et al. [14], where gold electrodes were previously applied to the sample. A strongly non-linear response of the  $I-V$  characteristic is observed whatever measurements are carried out through a grain boundary or in the grain. This behavior can be explained either by the intrinsic response of the CCTO or from a non-ohmic contact between the metallic tip and the specimen wherever it is contacted (inside a grain or between grain boundaries).

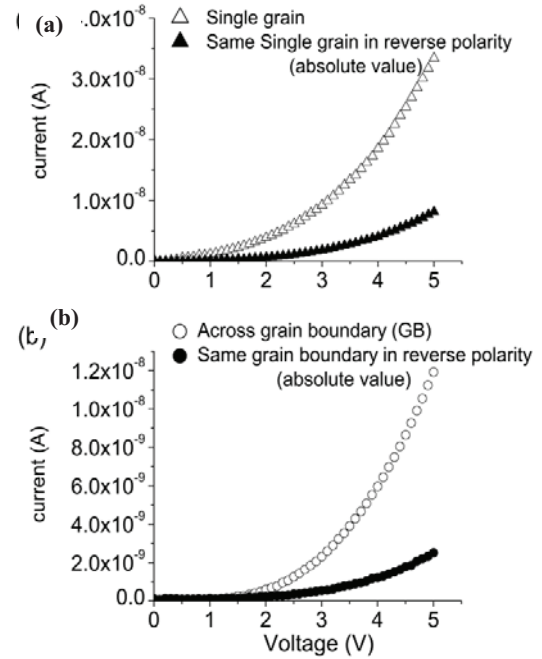


Fig. 11  $I-V$  measurements characteristics (a) within grains and (b) across individual grain boundaries at  $25$  °C.

Surprisingly, it is also observed, as in the case in bulk materials, a non-symmetrical response on both grain and across grain boundaries that is dependent on the polarity applied with the sourcemeter (Fig. 11). Care was taken in order to avoid moving tips in between the measurements, and polarity is changed by inverting wires connected to the voltage source. This result means that the total current passing in one direction in the measurement setup (probe contact, then ceramic) is different from the one in the opposite direction.

One possible explanation of all these measurements could be that the contact between an electrode and the material whatever the scale level, can be considered as non-ohmic. The material could behave in that case as a MIS (metal-insulator-semiconductor) structure and the IBLC model is no more necessary to explain the different behaviors.

#### 4. Conclusions

The use of characterization techniques that are complementary to impedance spectroscopy measurements, allowed to identify a non-symmetrical response relative to the applied voltage, in both bulk ceramic materials, and at the local level.

I-t measurements performed at different voltage levels, showed a high current related to the strong non-linear  $I$ - $V$  response of CCTO. When the applied voltage step is higher than 2 V, absorption and desorption currents are no longer super-imposable in the CCTO ceramics.  $I$ - $t$  measurement should complete, in future works, the panel of electrical characterization applied to this type of materials.

In regards to the  $I$ - $V$  responses, for bulk ceramics, or for the grains and grain boundaries, the interfaces contact between the sample and the metal electrode seems to be the main key to explain the non-linear and non-symmetrical observed behaviors and a model like MIS structure can be a plausible explanation of these characteristics.

Further investigations concerning the interfaces CCTO/electrodes can lead to a better understanding of the non-linear and non-symmetrical  $I$ - $V$  characteristics.

The current models describing interfacial polarization, either electrode/surface polarization or IBLC do not take into account the non-symmetrical phenomena, meaning that a more comprehensible model is yet to be built.

## References

- [1] M.A. Subramanian, D. Li, N. Duan, B.A. Reisner, A.W. Sleight, High dielectric constant in  $\text{ACu}_3\text{Ti}_4\text{O}_{12}$  and  $\text{ACu}_3\text{Ti}_3\text{FeO}_{12}$  phases, *Journal of Solid State Chemistry* 151 (2) (2000) 323-325.
- [2] A. Onodera, M. Takesada, K. Kawatani, S. Hiramatsu, Dielectric properties and phase transition in  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  at high temperatures, *Japanese Journal of Applied Physics* 47 (9) (2008) 7753-7756.
- [3] S. Guillemet-Fritsch, T. Lebey, M. Boulos, B. Durand, Dielectric properties of  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  based multiphased ceramics, *Journal of the European Ceramic Society* 26 (7) (2006) 1245-1257.
- [4] S. Krohns, P. Lunkenheimer, S. Ebbinghaus, A. Loidl, Broadband dielectric spectroscopy on single-crystalline and ceramic  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$ , *Applied Physics Letters* 91 (2) (2007) 022910-1-022910-3.
- [5] G. Deng, Z. He, P. Muralt, Physical aspects of colossal dielectric constant material  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  thin films, *Journal of Applied Physics* 105 (8) (2009) 084106.
- [6] L. He, J. Neaton, M.H. Cohen, D. Vanderbilt, C.C. Homes, First-principles study of the structure and lattice dielectric response of  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$ , *Physical Review B* 65 (21) (2002) 214112-1-214112-11.
- [7] P. Lunkenheimer, V. Bobnar, A. Pronin, A. Ritus, A. Volkov, A. Loidl, Origin of apparent colossal dielectric constants, *Physical Review B* 66 (5) (2002) 052105-1-052105-4.
- [8] G. Deng, T. Yamada, P. Muralt, Evidence for the existence of a metal-insulator-semiconductor junction at the electrode interfaces of  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  thin film capacitors, *Applied Physics Letters* 91 (20) (2007) 202903-1-202903-3.
- [9] D.C. Sinclair, T.B. Adams, F.D. Morrison, A.R. West,  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  one-step internal barrier layer capacitor, *Applied Physics Letters* 80 (12) (2002) 2153-2155.
- [10] J. Sebald, S. Krohns, P. Lunkenheimer, S. Ebbinghaus, S. Riegg, A. Reller, et al., Colossal dielectric constants: A common phenomenon in  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  related materials, *Solid State Communications* 150 (17) (2010) 857-860.
- [11] W.X. Yuan, Impedance and electric modulus approaches to investigate four origins of giant dielectric constant in  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  ceramics, *Solid State Sciences* 14 (3) (2012) 330-334.
- [12] T.B. Adams, D.C. Sinclair, A.R. West, Giant barrier layer capacitance effects in  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  ceramics, *Advanced Materials* 14 (18) (2002) 1321-1323.
- [13] J. Yang, M. Shen, L. Fang, The electrode/sample contact effects on the dielectric properties of the  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  ceramic, *Materials Letters* 59 (29-30) (2005) 3990-3993.
- [14] S.Y. Chung, I.D. Kim, S.J.L. Kang, Strong nonlinear current-voltage behaviour in perovskite-derivative calcium copper titanate, *Nature Materials* 3 (2004) 774-778.
- [15] P. Fiorenza, R. Lo Nigro, C. Bongiorno, V. Raineri, M.C. Ferarrelli, D.C. Sinclair, et al., Localized electrical characterization of the giant permittivity effect in  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  ceramics, *Applied Physics Letters* 92 (18) (2008) 182907-1-182907-3.
- [16] S.Y. Chung, Comment on: Origin of giant dielectric response in nonferroelectric  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$ : Inhomogeneous conduction nature probed by atomic force microscopy, *Chemistry of Materials* 20 (19) (2008) 6284-6285.
- [17] D. Fu, H. Taniguchi, T. Taniyama, M. Itoh, S.Y. Koshihara, Origin of giant dielectric response in nonferroelectric  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$ : Inhomogeneous conduction nature probed by atomic force microscopy, *Chemistry of Materials* 20 (5) (2008) 1694-1698.
- [18] Z. Valdez-Nava, S. Dinculescu, T. Lebey, Non-symmetrical electric response in  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  and  $\text{La}_{0.05}\text{Ba}_{0.95}\text{TiO}_3$ -SPS materials, *Journal of Physics D: Applied Physics* 43 (38) (2010) 385401-1-385401-6.
- [19] L. Marchin, S. Guillemet-Fritsch, B. Durand, A. Levchenko, A. Navrotsky, T. Lebey, Grain



- growth-controlled giant permittivity in soft chemistry  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  ceramics, *Journal of the American Ceramic Society* 91 (2) (2008) 485-489.
- [20] S. Krohns, P. Lunkenheimer, C. Kant, A.V. Pronin, H.B. Brom, A.A. Nugroho, et al., Colossal dielectric constant up to gigahertz at room temperature, *Applied Physics Letters* 94 (2009) 122903-1-122903-3.
- [21] Z. Valdez-Nava, S. Guillemet-Fritsch, C. Tenailleau, T. Lebey, B. Durand, J.Y. Chane-Ching, Colossal dielectric permittivity of  $\text{BaTiO}_3$ -based nanocrystalline ceramics sintered by spark plasma sintering, *Journal of Electroceramics* 22 (1-3) (2009) 238-244.
- [22] W. Li, R.W. Schwartz, Derivation and application of an empirical formula to describe interfacial relaxation effects in inhomogeneous materials, *Journal of the American Ceramic Society* 90 (11) (2007) 3536-3540.
- [23] T.B. Adams, D.C. Sinclair, A.R. West, Characterization of grain boundary impedances in fine- and coarse-grained  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$  ceramics, *Physical Review. B* 73 (9) (2006) 094124-1-094124-9.
- [24] G. Deng, P. Muralt, On origin and intrinsic electrical properties of the colossal dielectric constant state in  $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$ , *IOP Conf. Series: Materials Science and Engineering* 8 (2010) 012016-1-012016-4.