

Article

Improvement of the Electrical Performance of Outdoor Porcelain Insulators by Utilization of a Novel Nano-TiO₂ Coating for Application in Railway Electrification Systems

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Abstract: The present study aimed to develop the electrical performance of outdoor insulators using a nano-TiO₂ coating for railway electrification systems. The prototype design of porcelain insulators with normal coatings and using a nano-TiO₂ coating is based on IEC 60815-1. The first test was performed to measure the low-frequency flashover AC voltage under both dry and wet conditions. In addition, the other test was conducted to measure the lightning impulse critical-flashover voltage at positive and negative polarity under dry-normal and wet-contaminated conditions. X-ray diffraction (X-RD) and Scanning electron microscopy (SEM) were used to examine the micro surface and show that the nano-TiO₂ coating was adhered to the surface of the outdoor porcelain insulator and exists in an amorphous state. Additionally, it was observed and discovered that scattered nano-TiO₂ strengthens the glassy matrix and creates a sturdy barrier that causes flashover voltage to be reduced under conditions of high dielectric strength. Nanostructured ceramic formulations outperform ordinary porcelain in terms of breakdown voltage strength, particularly for the insulators' low-frequency flashover performances under dry and wet test conditions. However, a significant change in the lightning impulse critical-flashover voltage characteristics is observed and is not much better when adding the nano-TiO₂ coating to the porcelain insulators.

Keywords: railway electrification system; outdoor porcelain insulators; low-frequency flashover AC voltage; lightning impulse critical-flashover voltage



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1. Introduction

An important component of any electrical system is insulation. This includes gas insulation, such as SF₆, and liquid insulation, such as transformer oil or mineral oil, and solid insulation in transformers or cables, such as kraft paper or pressboard. In addition, there are important insulators that are used on the outside, such as the high-voltage porcelain insulators on the outside of transformer bushings, cutouts, arresters, line posts, etc. [1]. To last for years, they must meet strict electrical, mechanical, and chemical requirements, such as not breaking down when exposed to UV light or when in a dirty environment [2–4]. Although polymeric insulators have just lately been available, porcelain and glass have been used as outside insulators in electric power systems for over 160 years [5–8]. In the past, triaxial porcelain compositions of alumina, silica, clay, and feldspar were used to create ceramic insulators [1,9]. Insulators are one of the essential elements of railway electrification systems; however, the popular viewpoint does not give this essential component much attention. Economic considerations must be made in

addition to the strict criteria for availability and dependability, which is why this topic is given a lot of attention [10]. The dependability of the performance of insulators, which are widely used in transmission power systems, is primarily influenced by environmental pollution. Composite insulators have been used for transmission lines for a long time because they work well even in dirty environments [11]. As compared to other ceramic materials, outdoor porcelain insulators are the subject of little research today that deals with nanotechnology applications. Nevertheless, it has been said that they have produced some very intriguing and encouraging outcomes. Porcelain is a polycrystalline ceramic created by combining and burning clays, feldspathic minerals, and silica or alumina, and may be characterised as a conventional ceramic formed by heating raw materials based on clay [6,12]. In 2010, Zhuang et al. [13] investigated a novel use of nanoscale titanium dioxide (TiO_2) for improving the performance of the porcelain insulator in contaminated settings. The researchers found that applying $n\text{-TiO}_2$ films to the porcelain insulator's surface maintains and even enhances the electrical characteristics of the AC wet flashover voltage (6%). To enhance the anti-icing performance of ceramic insulators utilised in China's colder regions, Gou et al. created three novel experimental formulations of polymeric nanocoatings [14]. In order to enhance the mechanical and electrical performance of siliceous porcelain insulators, Contreras investigated a novel nanotechnology concept in 2014 [1].

At present, the rail system is of interest in the transportation system of Thailand. Therefore, it is important to research and develop the design of the components used in the rail system. In particular, the insulation in the railway electrification systems is one of the details of the overhead line electrification system for the railway, as can be seen in Figure 1. In addition, Thailand's rail transport system uses an overhead power supply system to operate. Therefore, it is necessary to instal insulating materials, including outdoor porcelain insulators. There is very little research about the design and build of the porcelain insulator used in the electrification system for the railway in Thailand. Furthermore, most of these studies were about testing the durability of insulators, which included electrical tests.

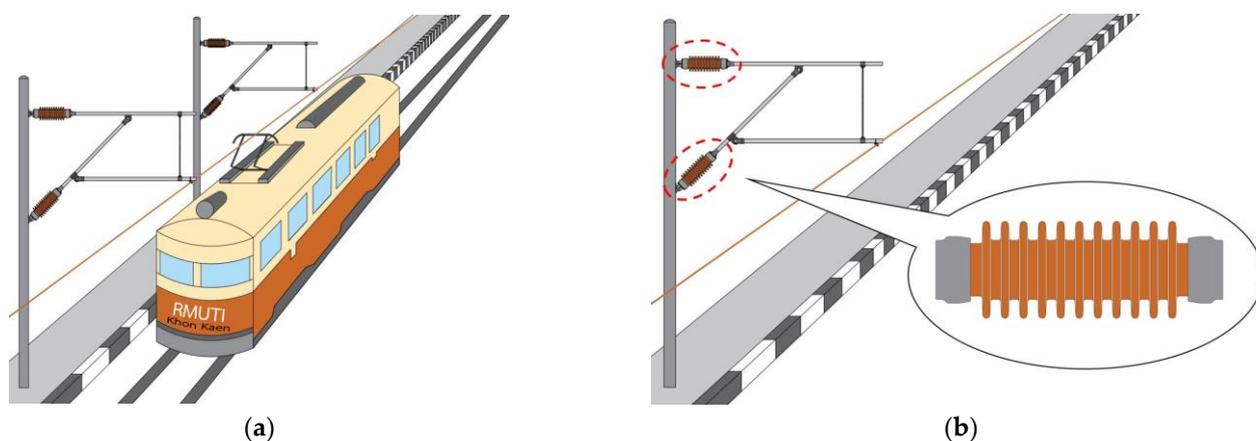


Figure 1. Railway electrification system: (a) Overview of railway electrification system; (b) Design of overhead line insulation equipment as porcelain insulator.

This work was performed to improve the quality of the outdoor porcelain insulators used in the railway system and make it possible for electric trains to run on Thailand's railway system. The porcelain insulator used in this railway system is extremely important because there is no manufacturer in Thailand, as they are primarily interested in producing insulators for the alternating current distribution system. As a result, the goal of this research is to develop the performance of outdoor insulators used in the railway's overhead line electrification system. The performance-developed outdoor insulators of overhead line insulation equipment for railway electrification systems that use a nano- TiO_2 coating would be the quality insulator that passes the standard electrical properties test.

2. Materials and Methods

2.1. The Design for Outdoor Insulators on Railway Electrification Systems

The prototype design of porcelain insulators based on international standards and the outdoor insulators on railway electrification systems used in Japanese and Spanish railway systems was studied, which included the equations used in the design of the leakage distance of insulators used in rail systems. Furthermore, the reference to the design of porcelain insulators, the insulator profile parameters as defined in IEC 60815-1 [15], the design considers the level of pollution that is used. Moreover, this research will design insulators that can be used at pollution levels of 1 and 2 based on IEC 60815-1. To determine the leakage distance of insulators, it is necessary to calculate the profile factor (PF) and the creepage factor (CF), which are important in the design of the fin length (measured from the core) and the distance between the fins, which includes the creepage distance and arcing distance of the insulators. Meanwhile, the values calculated from Equations (1) and (2) are evaluated to determine the suitability of insulating insulators for the design or application area. Previously, the sizes were as shown in Figure 2.

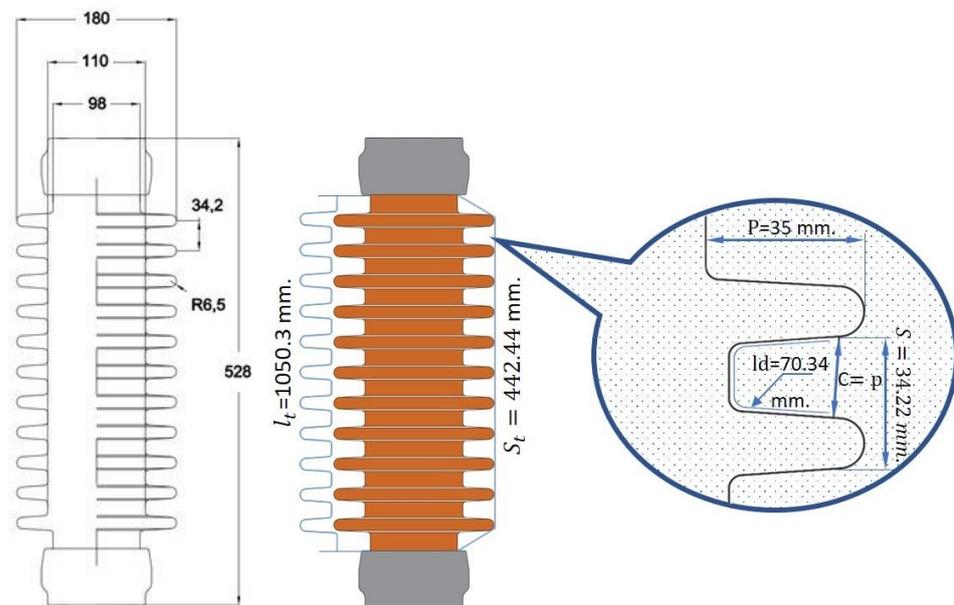


Figure 2. Schematic diagram of the main nanotechnology design concepts for outdoor insulators.

Profile Factor (P.F.)

$$P.F. = \frac{2P + S}{l_d} \quad (1)$$

when

P.F. > 0.8 for the pollution levels of 1 and 2 (light to medium pollution areas)

P.F. > 0.7 for the pollution levels of 3 and 4 (heavy to very heavy pollution areas)

Creepage Factor (C.F.)

$$C.F. = \frac{l_t}{S_t} \quad (2)$$

when

C.F. ≤ 3.5 for the level of pollution 1 and 2 (light to medium pollution areas)

C.F. ≤ 4.0 for the level of pollution 3 and 4 (heavy to very heavy pollution areas)

The outdoor insulator profile parameters in this work, as defined in IEC60815-1, are as follows:

P: shed projection—the maximum shed overhang (35 mm).

S : shed spacing—the vertical distance between two points that are similar on successive sheds. (34.22 mm).

l_d : Measured between the two places that constitute d is the creepage distance (70.34 mm).

l_t : the insulator's overall creepage distance (1050.30 mm).

S_t : the arcing distance of the insulator (442.44 mm).

The design variables were substituted into Equation (1) for P.F. and Equation (2) for C.F. So, the profile factor (P.F.) is equal to 1.482, which is more than 0.8 for the level of pollution in 1 and 2 (light to medium pollution areas). Moreover, the creepage factor (C.F.) is equal to 2.374, which is less than 3.5 for the level of pollution in 1 and 2 (light to medium pollution areas).

Therefore, the prototype outdoor insulators designed in this research were at pollution levels 1 and 2. It is standard practice on IEC60815.

2.2. Nanotechnology Concepts for Outdoor Insulators

This work presents new nanostructured porcelain composition coats for improving the electrical performance of outdoor porcelain insulators by using a novel application of nano-TiO₂ coatings, including evaluating the effect of nano-TiO₂ on the properties of conventional siliceous electrical porcelain for use in railway electrification systems. All the unprocessed nanomaterials employed in this investigation had an average particle size of ~40 nm. Additionally, they had a spherical form, and their analytical reagent quality was measured by TESCAN, Model: MIRA3, as seen in Figure 3.

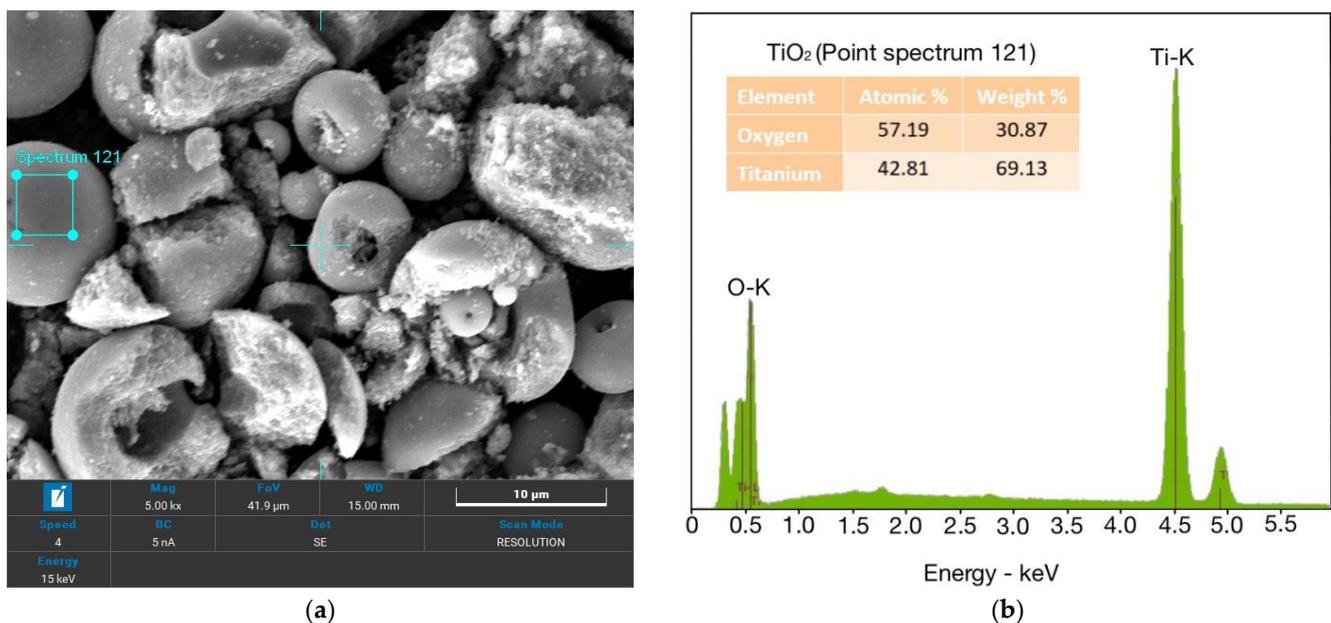


Figure 3. SEM nano-TiO₂ are as follows: (a) The morphology of nano-TiO₂; (b) The electron dispersive X-ray spectroscopy (EDS) result of nano-TiO₂ used in this work.

The shape of nano-TiO₂ is also shown in Figure 3a, where a quasi-spherical form is often seen. There could be some nano-TiO₂ agglomerates present since no dispersant was utilised. Electron dispersive X-ray spectroscopy (EDS) was used to confirm the chemical composition of nano-TiO₂, which is shown in Figure 3b. The properties of the nanoparticles employed in this work are listed in Table 1 [16]. Detailed properties of nano-TiO₂ are specified by the manufacturer [16]. Titanium dioxide (titania) nanopowder, anatase. Specification: 99.9%; anatase; primary particle size ~40 nm (from SEM and BET surface area); BET multi-point specific surface area (SSA) >40 m²/g; spray-dry agglomerated to ~50 microns for easy handling.

Table 1. The main properties of TiO₂ nanomaterial used to coat porcelain insulators are listed.

Characteristics	Specification
Parameters	Titanium dioxide (TiO ₂)
Average particle size	~40 nm
Purity	99.9%
Specific surface area	40 m ² /g
Structure	anatase
Colour	white powder
Density	3.89 g/cm ³

The key ideas in this study for the outdoor porcelain insulators used in the railway system represent a novel approach to improving the final characteristics of outdoor high-voltage insulators. In addition, the porcelain raw materials utilised in this study were of industrial quality and were quartz, kaolinite clay, and sodium feldspar, which are all employed in the production of outdoor insulators according to the expertise of Asian Insulators Public Company Limited. Figure 4 shows that the main use of nanotechnology in this outdoor insulator by nano-TiO₂ coating is to improve efficiency, mostly by making the electrical performance better.

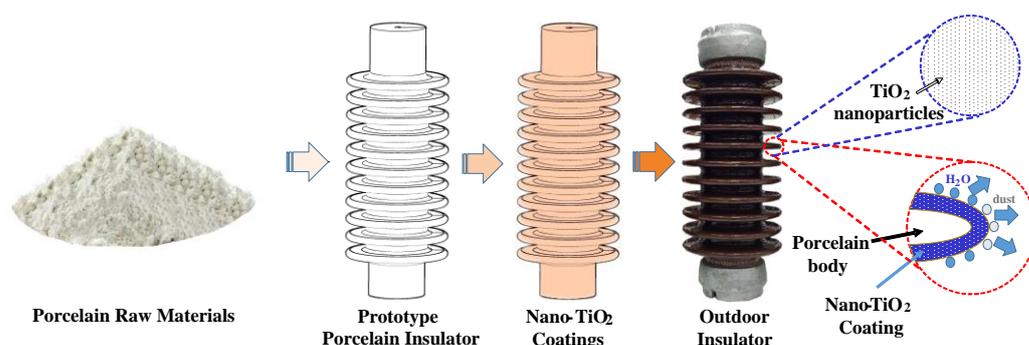
**Figure 4.** Schematic of the design concepts for nano-TiO₂ coatings of outdoor insulators in this study.

Figure 4 illustrates the design idea for the aforementioned procedure. The link between a material's resistivity, length, and the area is shown by the well-known Equation (3) [17].

$$R = \rho \frac{L}{A} \quad (3)$$

where R stands for resistance, ρ for resistivity, L for the distance that charges travel before leaving the material, and A for the cross-sectional area of the material that faces the direction of the current. The R will decrease if we shorten L , raise A , or decrease ρ , as shown by Equation (3).

For the purposes of this study, we will refer to ρ as the resistivity of the nano-TiO₂ coating, L as the thickness of the nano-TiO₂ film, and A as the surface area of the graphite felt used for the nano-TiO₂ film deposition. Thus, lowering the size and L of the nano-TiO₂ film is one of the development paths that might be explored in this study. In this work, a nano-TiO₂ coating was formed on the outside insulators to shorten L in order to decrease the resistance (R) of the coating. On the other hand, a two-step procedure was employed to create a nano-TiO₂ coating, which included first generating an amorphous film at a low temperature, and then increasing the temperature to crystallise the film. It is noted that a two-step technique may be used to create a nano-TiO₂ coating [18–20]. Since a large number of grain boundary defects in the nano-TiO₂ coating, the resistivity (ρ) of the nano-TiO₂ coating can be effectively reduced. Besides, the nano-TiO₂ coating structure can achieve low resistance.

2.3. Surface Modification of Outdoor Insulators

A thin nano-TiO₂ layer was applied to the surface of outdoor insulators via the development of an insulator nano-TiO₂ coating method for surface modification. The process includes six steps. The nanoparticles are mixed with deionized water. The nanoparticles must be weighed on a digital balance, and the particle content required is 0.1% wt of deionized water. TiO₂ nanoparticles (nano-TiO₂) combined with deionized water and a surfactant. The surfactants are organic compounds, which consist of two parts: the hydrophilic group and the hydrophobic group. Two different phases can come together. When a small amount of surfactant is added, two phases can be combined to form a single phase. In this research, a surfactant type, TERGITOL™ NP-6 Surfactant (Nonylphenol Ethoxylate) [21], which has properties as a non-ionic surfactant, was selected to be used as an ingredient in the preparation of nano-TiO₂ coatings for outdoor porcelain insulators. Then, it is placed in the stirrer for 30 min at 60 °C, or until it diffuses, and put in the high-frequency ultrasonicator at 60 °C for 1 h to ensure the homogeneity of the liquids, as shown in Figure 5. The step, as shown in Figure 5, begins with weighing the normal coatings and nano-TiO₂ coatings at the prescribed amount (a weight ratio of 2:1 between the normal coatings and nano-TiO₂ coatings). The next process involves mixing two substances, including measuring, and controlling the coating viscosity at 30–45 Cps. and waiting for the next process. After that, the nano-TiO₂ coatings are put into a chamber containing the coatings. The formed prototype insulator is then put on an automatic coating machine to process the coating. Lastly, the prototype insulator is put into a temperature-controlled kiln under control at 1300 °C for 60 h.

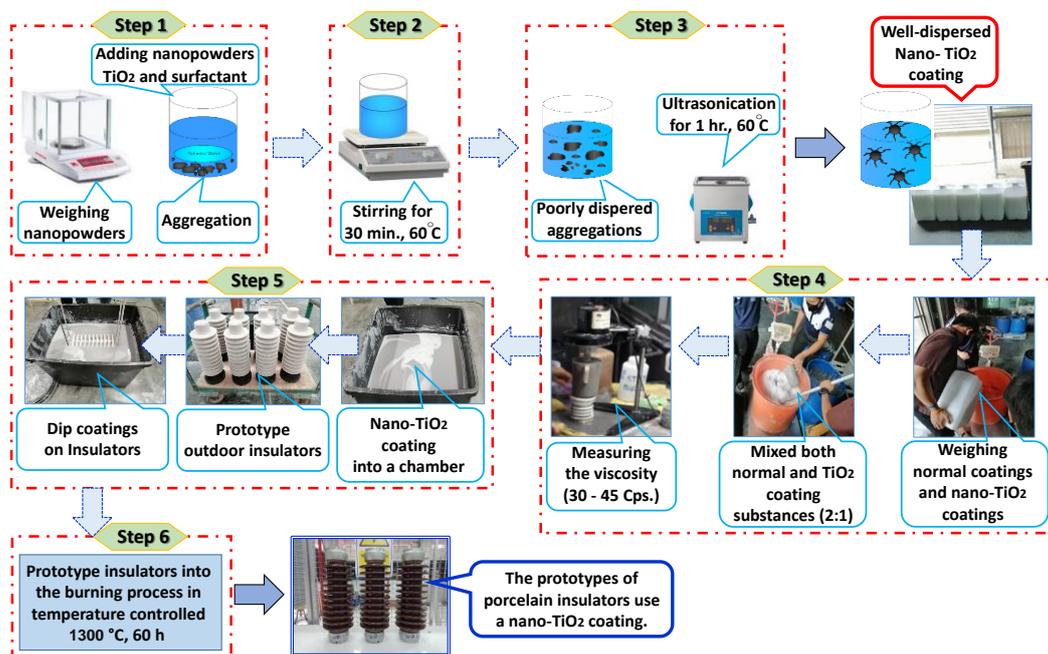


Figure 5. Preparation of nano-TiO₂ coating of outdoor insulators in this study.

3. Experimental Descriptions

3.1. Performances of Standard Porcelain Insulators in Low-Frequency Flashover in Dry and Wet Test Conditions

When referring to low frequency, as defined in these standards, anything between 15 and 100 cycles per second is considered to be low frequency. The low-frequency testing described in these standards may be performed at any commercial frequency that is currently in use within this range. In Figure 6, a clean, dry test specimen voltage application's root-mean-square (rms) voltage that results in a prolonged disruptive discharge through the air between electrodes is known as a low-frequency dry-flashover voltage. Approximately

75% of the anticipated average dry-flashover value may be rapidly reached by rapidly increasing the first applied voltage. The rate of voltage rise must remain constant after achieving 75% of the flashover value in order for there to be no time delay of more than 30 s or less than 5 s before the flashover occurs. An arithmetic mean, given in rms volts, of at least five different flashovers collected in a row constitutes the dry-flashover voltage value for a specimen. There must be between 15 s and 5 min of time between each flashover.

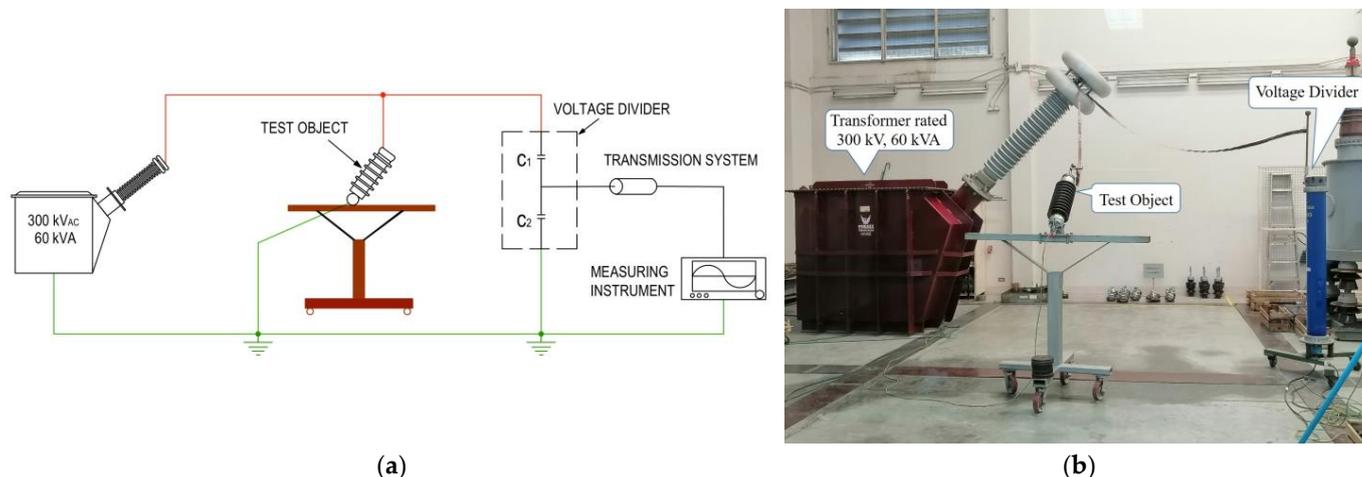


Figure 6. The experimental circuit of low-frequency flashover AC voltage under dry test. (a) The circuit diagram on low-frequency dry-flashover test; (b) The test circuit set-up on low-frequency dry-flashover test.

The low-frequency wet-flashover voltage is the root-mean-square (rms) voltage that causes a long, disruptive discharge between the electrodes of a clean, wet test specimen. The area where the insulator is being tested must have enough spray nozzles of the kind illustrated in Figure 7 or an equivalent supply of artificial precipitation that is fairly uniform. Within the limits of what is practically possible, the spray must be directed such that it strikes the insulator downward at a 45-degree angle from the vertical and parallel to the vertical plane across its axis. Large specimens, such as the outdoor porcelain insulators, must have spray that is 45 degrees off vertical in the middle and as near to this angle at the ends as feasible to provide adequate precipitation uniformity over the length of the specimen. At the specimen, the standard average rate of precipitation must be 0.2 inches (5.08 millimetres) per minute when measured. The low-frequency flashover AC voltage testing apparatus is shown in Figures 6 and 7 for both dry and wet experiments, respectively. Phenix Technologies' variants for the frequency output device are rated at 300 kV, 60 kVA, respectively, based on ANSI/NEMA C29.7-2015 [22], TIS 2623-2560 (2017) [23], and IEEE-4 (2013) [24].

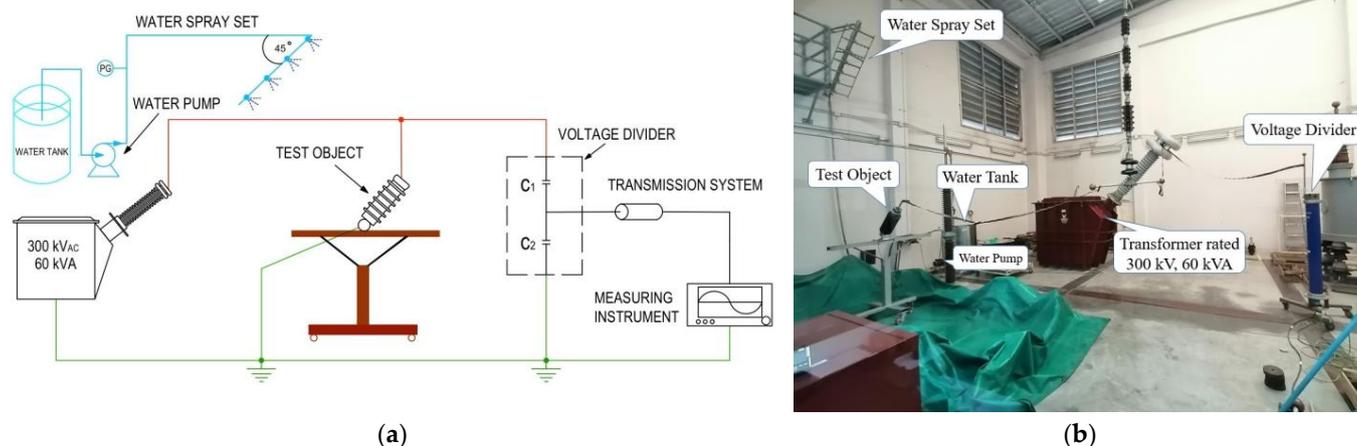


Figure 7. The experimental circuit of low-frequency flashover AC voltage under wet test. (a) The circuit diagram on low-frequency wet-flashover test; (b) The test circuit set-up on low-frequency wet-flashover test.

3.2. The Lightning Impulse Critical-Flashover Voltage Performances of Standard Porcelain Insulators under Normal and Contaminated Test Conditions

A test specimen's impulse critical-flashover voltage is equal to the wave's peak value in 50% of applications. A lightning impulse critical-flashover voltage test in polluted circumstances was used in this study to assess the performance of prototype outdoor porcelain insulators. Normally, impulse-flashover experiments are only carried out in dry environments. There must be 1.2/50 μ s of time between each impulse voltage wave. Applying impulses to the specimen of varied intensities and monitoring the peak voltage of the impulse that, in about 50% of applications, will result in flashover allows for the determination of the critical impulse flashover voltage value. Impulses must be applied in stages of rising or decreasing voltage such that at the step with the greatest voltage, flashover occurs on every impulse, and at the step with the lowest voltage, there is no flashover. At least three impulses must be applied at each voltage step, and the difference in voltage between steps must not be more than 5% of the anticipated critical impulse flashover value. When the voltage range between the highest and lowest step is quite large, it is possible to calculate the crucial impulse flashover voltage by plotting an average curve of crest voltage vs flashover frequency.

The experimental setup for testing the critical-flashover voltage for a lightning impulse in both dry and polluted conditions is shown in Figures 8 and 9. The impulse voltage generator, whose rated voltage is 400 kV and 40 kJ under both normal and contaminated circumstances based on TIS 1077-2535 (1992) [25], TIS 2623-2560 (2017), and IEC 60507 (2013) [26], is used to acquire test power. The primary capacitor is charged to the desired voltage. The circuit is then automatically initiated by the spark gap to provide a positive polarity lightning discharge voltage of 1.2/50 μ s. The experiment uses the up-and-down test methodology. A test was conducted in order to determine how effectively conventional porcelain insulators can tolerate impulses at both positive and negative polarities and in unclean circumstances. Furthermore, the surface of the specimens is polluted using test techniques on pollution flashover the suspension of sodium chloride and kieselguhr, which, respectively, imitate electric and inert materials. Based on the results of [27–29], the relationship between the critical flashover voltage representations of the electrical withstand value and the equivalent salt deposit density (ESDD) values is recommended as a contamination severity criterion for each case, as shown in Equation (4):

$$E_s = \lambda(ESDD)^{-\mu} \quad (4)$$

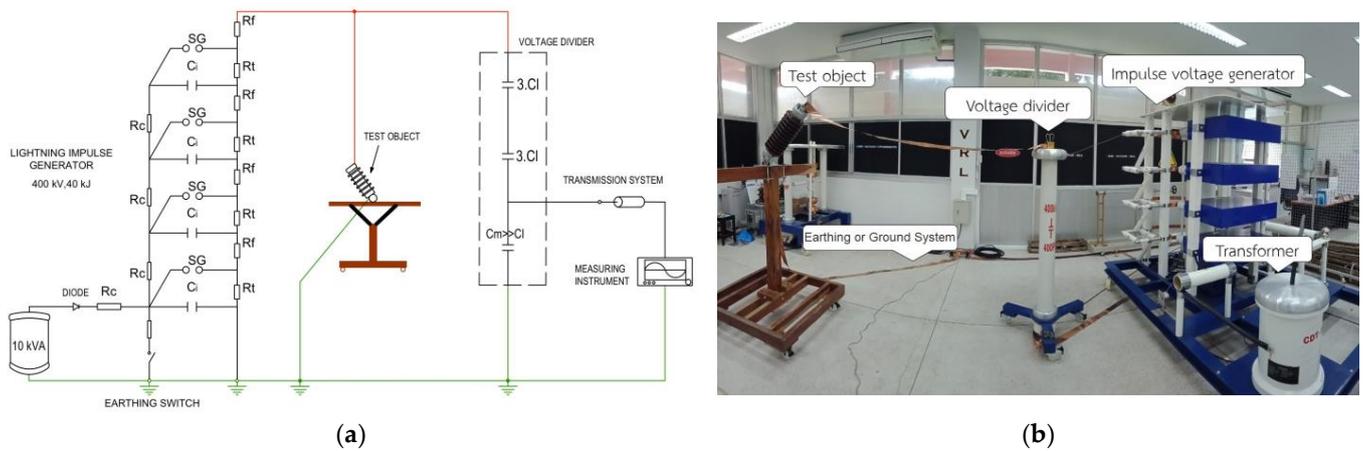


Figure 8. The experimental circuit of the lightning impulse critical-flashover voltage under normal condition test. (a) The circuit diagram on the lightning impulse critical-flashover voltage test; (b) The test circuit set-up on the lightning impulse critical-flashover voltage test.

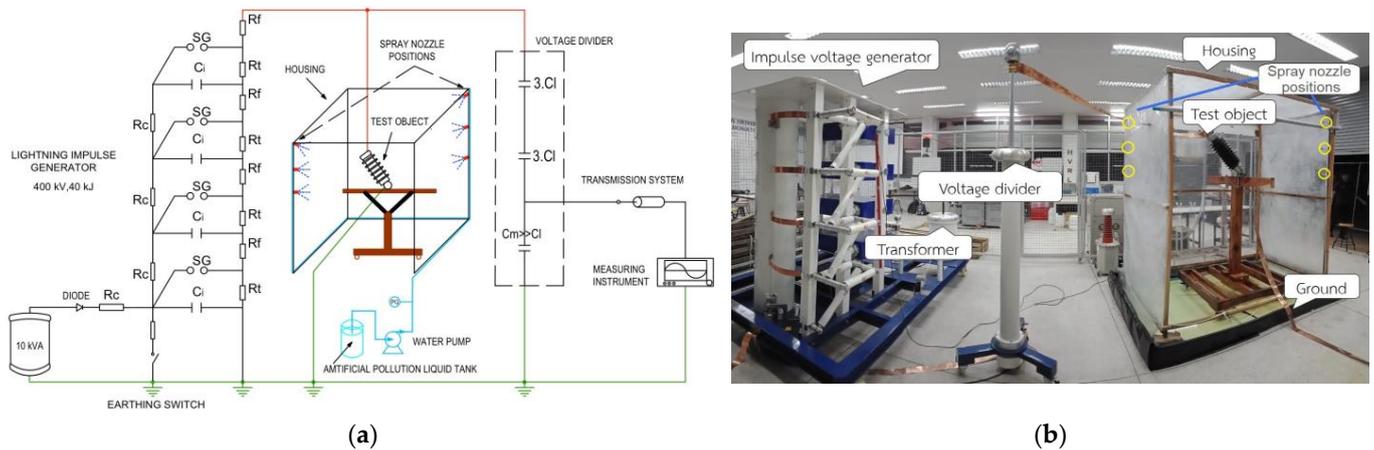


Figure 9. The experimental circuit of the lightning impulse critical-flashover voltage under contaminated condition test. (a) The circuit diagram on the lightning impulse critical-flashover voltage test; (b) The test circuit set-up on the lightning impulse critical-flashover voltage test.

When the electrical voltage is withstood (E_s), which is measured using experimental data from research and curve fitting, it is written as kV/cm and is related to the material and shape of the insulators. This research provides samples of contaminated liquid for testing under pollution conditions. By using the data for the contaminated liquid samples, values of ESDD can be calculated based on the procedure described in IEC 60507. The ESDD value obtained in this work is 0.4 mg/cm^2 .

For all experiments, the salt deposit density (SDD)/equivalent salt deposit density ratio is 1:6. At the top of the artificial climate chamber, the test specimen is mounted on a solid base made of non-conductive wood. Spray nozzles are used to spray the test specimen. A spray pipe that is located not far below the surface of the body of the housing carries the spray, and the spray mist input rate is $(0.05 \pm 0.01) \text{ kg}/(\text{h} \cdot \text{m}^3)$ based on IEEE 4 (2013), and the fog chamber's humidity is almost saturated to a wet condition [30]. Through exhausting and misting water, the mist chamber is cooled to match the ambient temperature. Once the insulator's surface is suitably moist, the sample is left to stand for 15 to 20 min before the test is begun [26].

4. Results and Discussion

4.1. The Dielectric Strength

4.1.1. Low-Frequency Flashover Performance in Dry and Wet Testing Conditions

The influence of nano-TiO₂ on the low-frequency flashover AC voltage under dry and wet tests of the prototype outdoor porcelain insulator sample is shown in Table 2. It was discovered that adding nano-TiO₂ encouraged an improvement in the dielectric capacity of regular porcelain.

Table 2. Test results of low-frequency flashover performances under dry and wet test condition (kV).

Test Conditions	Insulator Types	1st	2nd	3rd	4th	5th	Average	SD
Standard valve of condition test, kV (125 kV, based on ANSI/NEMA C29.7-2015)								
Dry	Normal coating	174.33	172.44	171.49	172.44	173.38	172.81	1.07
	Nano-TiO ₂ coating	176.21	174.30	175.26	176.21	174.30	174.28	0.95
Standard valve of condition test, kV (95 kV, based on ANSI/NEMA C29.7-2015)								
Wet	Normal coating	156.79	158.81	157.80	159.06	157.80	158.05	0.90
	Nano-TiO ₂ coating	159.90	160.41	158.59	159.40	160.42	159.74	0.77

SD = Standard deviation of the data on low-frequency flashover performances tests.

Figure 10a depicts dry test results with the nano-TiO₂ concentration used in this study (0.1 wt.%), with the breakdown voltage reaching an average of 174.28 kV at 0.1 wt.% nano-TiO₂ addition and 172.81 kV at 0 wt.% nano-TiO₂ addition (uncoated nano-TiO₂). Using a new way to use nano-TiO₂ coatings on samples made the electrical performance of outdoor porcelain insulators 0.85% better than with regular porcelain. The effect of the nano-TiO₂ on the low-frequency flashover AC voltage under wet tests of the porcelain insulator sample is depicted in Figure 10b. It was found that the addition of nano-TiO₂ impacts the electrical performance of the porcelain insulator. When nanostructured porcelain with a nano-TiO₂ coating was compared to regular porcelain (without a nano-TiO₂-coating), it was better by 1.06%. Due to the inclusion of nanoparticles, the sintered microstructure has changed, which may be the cause of this behavior. According to [31,32], a larger concentration of the vitreous phase has a negative impact on the dielectric breakdown because the crystalline/glassy phase ratio affects the electric characteristics of porcelain systems. Due to a lower proportion of glassy phase than normal porcelain, nanostructured porcelain exhibits greater breakdown voltage values. This behaviour may be made possible by the optimization of the electrical design of insulators. Without a doubt, adding nano-TiO₂ to the prototype outdoor porcelain insulator makes its electrical properties better.

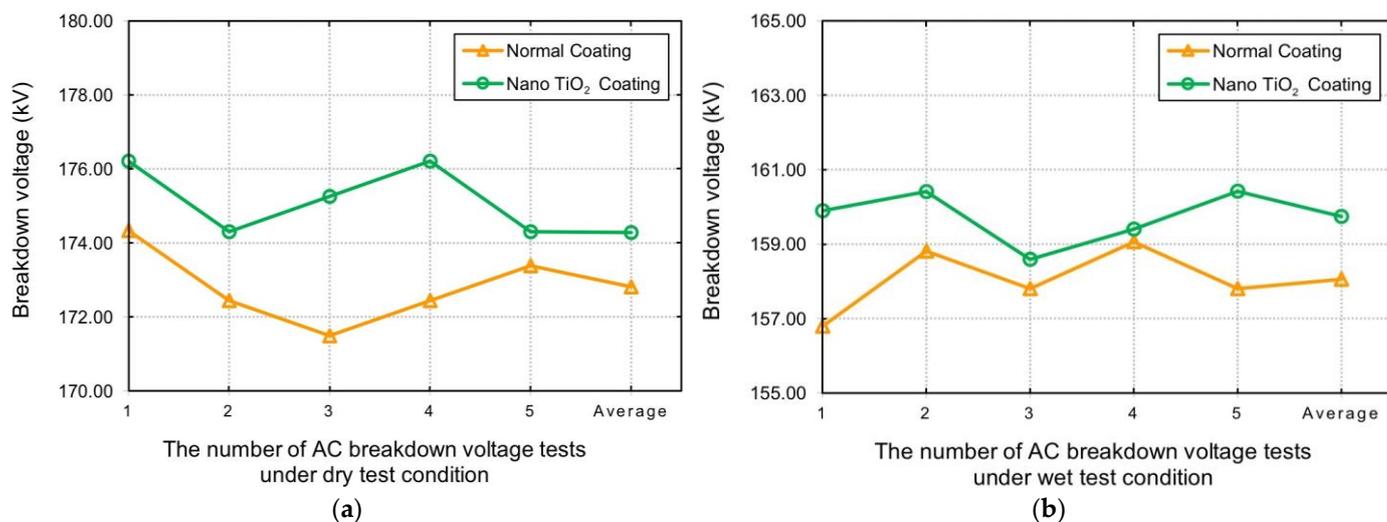


Figure 10. Test results of low-frequency flashover performances. (a) The low-frequency flashover performances under dry test condition; (b) The low-frequency flashover performances under wet test condition.

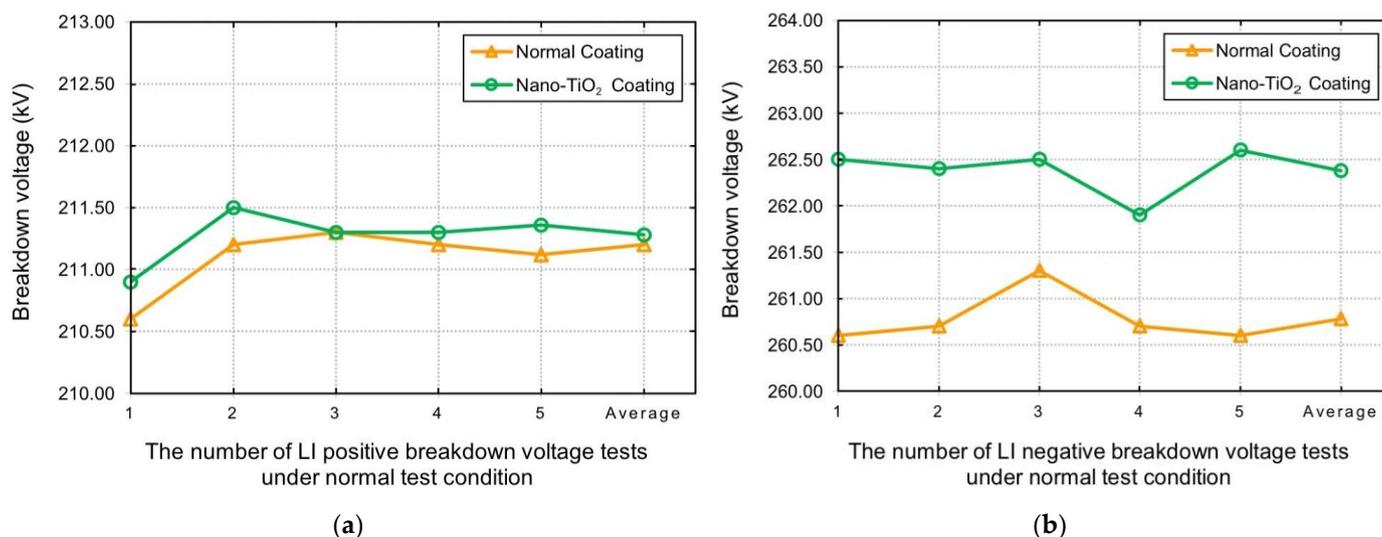
4.1.2. The Lightning Impulse Critical-Flashover Voltage Performance under Normal and Contaminated Test Conditions

A comparison of the critical impulse flashover voltage is illustrated in Table 3. In Figure 11, under normal test conditions, both positive and negative lightning impulses, the nano-TiO₂ coated insulators had a slight increase in dielectric strength compared to traditional coated insulators. In the same way, Figure 12 shows the critical impulse flashover voltage value under contaminated test conditions. When the nano-TiO₂ coating was added to the porcelain insulators, there was only a slight improvement in the changes under positive polarity testing, and in the negative polarity testing, there was no significant change in the characteristics of the lightning impulse critical-flashover voltage. The electrical performance test results are less than 2%, according to the results. It may be because the nano-TiO₂ content is so low (only 0.1 wt.%) that it does not show a clear effect on the electrical properties of the coated insulators with improved coating as much as it should. In the future, we may add more nano-TiO₂ to show higher electrical values. Additionally, it is well known that under both positive and negative lightning impulses, the distribution of the electric field on the insulator's surface is not uniform. The accumulation and growth of space charges would greatly distort the electric field across the air gap and reduce the flashover voltage [13,33]. Undoubtedly, one of the most effective ways to reduce the likelihood that charges will accumulate on the surface of an insulator is to decrease the surface resistivity [34]. Insulators are coated with nano-TiO₂ to make the insulator surface diffuse the electric field and thus increase the lightning impulse voltage. Contrarily, the surface conductance of the TiO₂-coated insulator is higher than that of the uncoated one, which might lead to an increase in leakage current and a decrease in flashover voltage. During testing, the U50% stays pretty much the same because these two opposing effects often cancel each other out [13].

Table 3. Test results of critical impulse flashover voltage value under normal and contaminated test conditions.

Test Conditions	Insulator Types	1st	2nd	3rd	4th	5th	Average	SD	
Test condition of critical impulse flashover voltage value, kV (210 kV, based on TIS. 1077-1992)									
Normal (dry)	LI Positive	Normal coating	210.6	211.2	211.3	211.2	211.3	211.12	0.27
		Nano-TiO ₂ coating	210.9	211.5	211.3	211.3	211.8	211.36	0.22
Test condition of critical impulse flashover voltage value, kV (260 kV, based on TIS. 1077-1992)									
LI Negative	Normal coating	260.6	260.7	261.3	260.7	260.6	260.78	0.29	
	Nano-TiO ₂ coating	262.5	262.4	262.5	261.9	262.6	262.30	0.27	
Comparison of critical-flashover voltage performance, kV									
Contaminated (wet)	LI Positive	Normal coating	147.1	148.2	147.4	149.2	148.4	148.06	0.83
		Nano-TiO ₂ coating	149.4	148.3	148.4	149.4	148.5	148.80	0.55
Comparison of critical-flashover voltage performance, kV									
LI Negative	Normal coating	185.5	184.9	185.6	185.2	183.7	184.98	0.76	
	Nano-TiO ₂ coating	185.4	184.0	183.7	183.9	183.6	184.12	0.73	

SD = Standard deviation of the data on lightning impulse (LI) tests.

**Figure 11.** Test results of critical impulse flashover voltage value under normal test conditions are as follows. (a) The lightning impulse critical-flashover voltage performance under positive polarity testing; (b) The lightning impulse critical-flashover voltage performance under negative polarity testing.

4.2. Phase Analysis

The prototype scale study's comparative XRD analysis of traditional porcelain systems and nanostructured porcelain systems in the sintered condition. Figure 13 shows the XRD patterns of prototype porcelain insulators with conventional coatings and nano-TiO₂ coatings measured by the Bruker D2 phaser at 30 kV, 10 mA. The results of this study revealed that the primary crystalline phases in the porcelain insulator prototype composition were quartz (SiO₂) and mullite (3Al₂O₃•2SiO₂). The primary crystalline phases of the nanostructured porcelain's diffraction pattern remained the same, and corundum, an unaltered form of alumina, was also present. Figure 13a shows alumina peaks in formulations with a high proportion of traditional porcelain insulators. Furthermore, as illustrated in Figure 13b, adding a TiO₂ layer to the surface of outdoor insulators improves mullite peak intensity. Additionally, the fraction of crystalline phases rise with the inclusion of nanoparticles. It

is important to note that the phase concentrations of all the samples match the published values for porcelain insulators [6,35,36]. The data for the prototype porcelain insulators with regular coatings and with a nano-TiO₂ coating were refined using the Rietveld method to produce crystallographic data, which was then used to further characterise the XRD findings. The lattice parameters, atomic coordinates, and atomic occupancies of manufactured materials may all be examined using the Rietveld refinement method, are displayed in Figure 14.

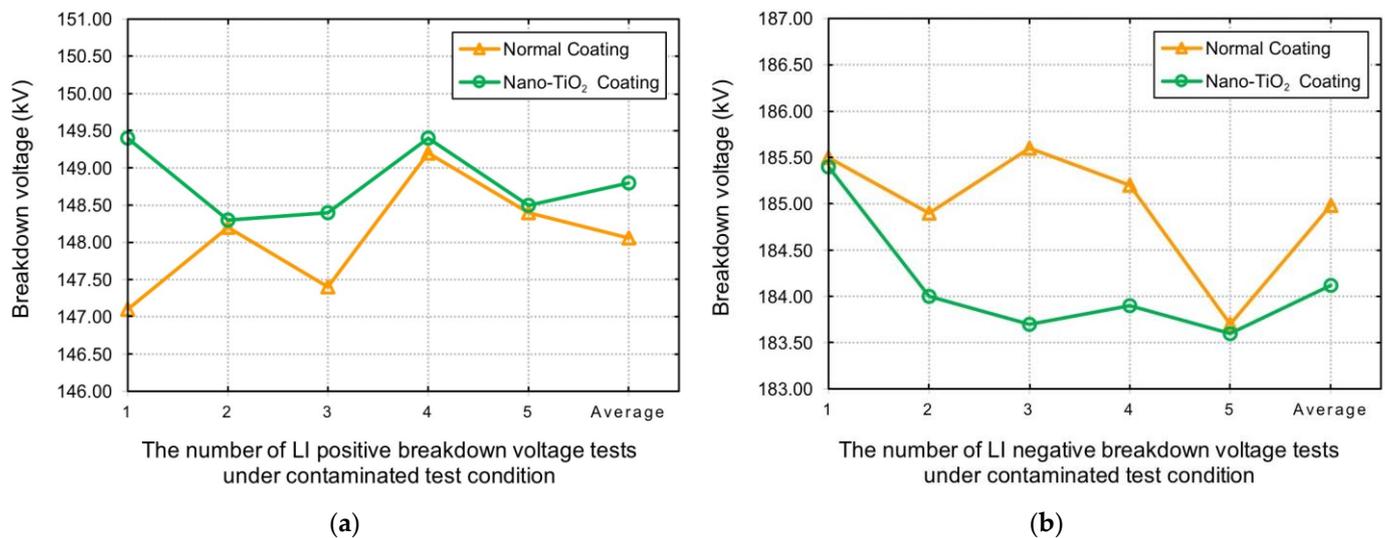


Figure 12. Test results of critical impulse flashover voltage value under contaminated test conditions are as follows. (a) The lightning impulse critical-flashover voltage performance under positive polarity testing; (b) The lightning impulse critical-flashover voltage performance under negative polarity testing.

The Rietveld refinement was carried out on porcelain insulators with normal coatings, and a nano-TiO₂ coating was used to analyse the crystal structure of the porcelain materials. Figure 14 shows the Rietveld refinement plot of the triaxial ceramic fired at 1300 °C. In this work, the observed XRD patterns and the structural model showed convergence with good weight profile R values (Rwp). The R values, Rietveld refinement, Residual values, or Rietveld discrepancy indices are all names for the numerical approach of evaluating the quality or goodness of fit, which is often expressed in terms of agreement indices [37]. The weighted profile R values are the easiest to understand because they come straight from the square root of the minimum amount, which is scaled using weighted intensities. Typical values of Rwp range from a few per cent for very accurate refinements to 20–30% for X-ray refinements, depending in part on the counting times used, the degree of preferred orientation, and the number of variable parameters [38,39]. The refinements were adequate. Indeed, the Rwp values were, in all cases, between 10 and 30%. Furthermore, in all samples, the estimated standard deviation of weight per cent derived from the estimated standard deviations of individual scale factors for the respective phases was below 1.5%. In particular, when considering the comparisons from Figures 14a and 14b, the graphs show the percentage of the substance content on all the samples of insulating coating. Moreover, it can be seen that when nano-TiO₂ coatings are applied to insulators, they result in higher percentages of mullite, and sillimanite compared to insulators using traditional coatings. The results obtained from Rietveld refinement are consistent with the XRD results shown in Figure 13.

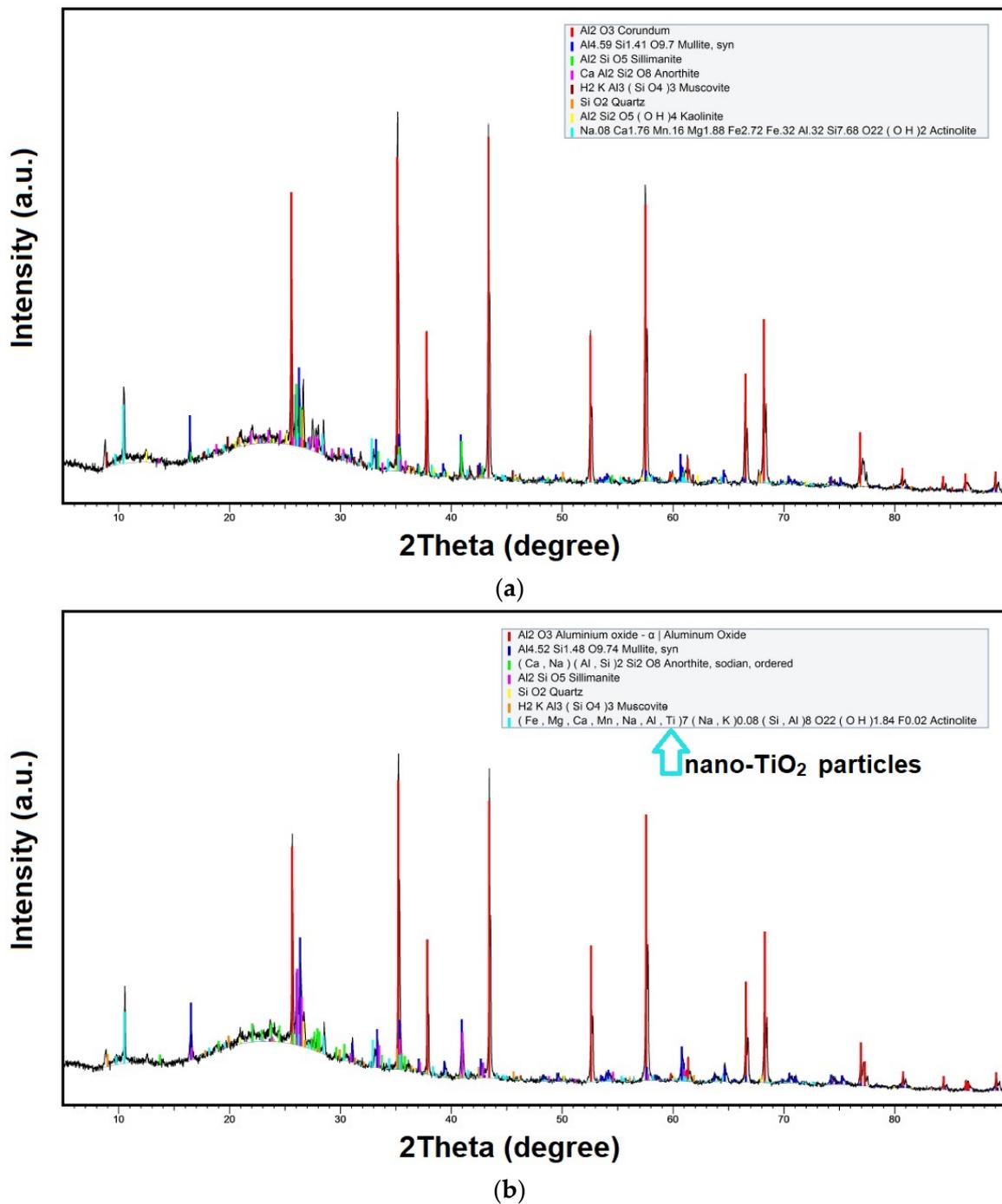


Figure 13. X-ray diffraction (X-RD) phase analysis. (a) X-RD pattern of a normal porcelain insulator prototype; (b) X-RD pattern of a nano-TiO₂ coated porcelain insulator prototype.

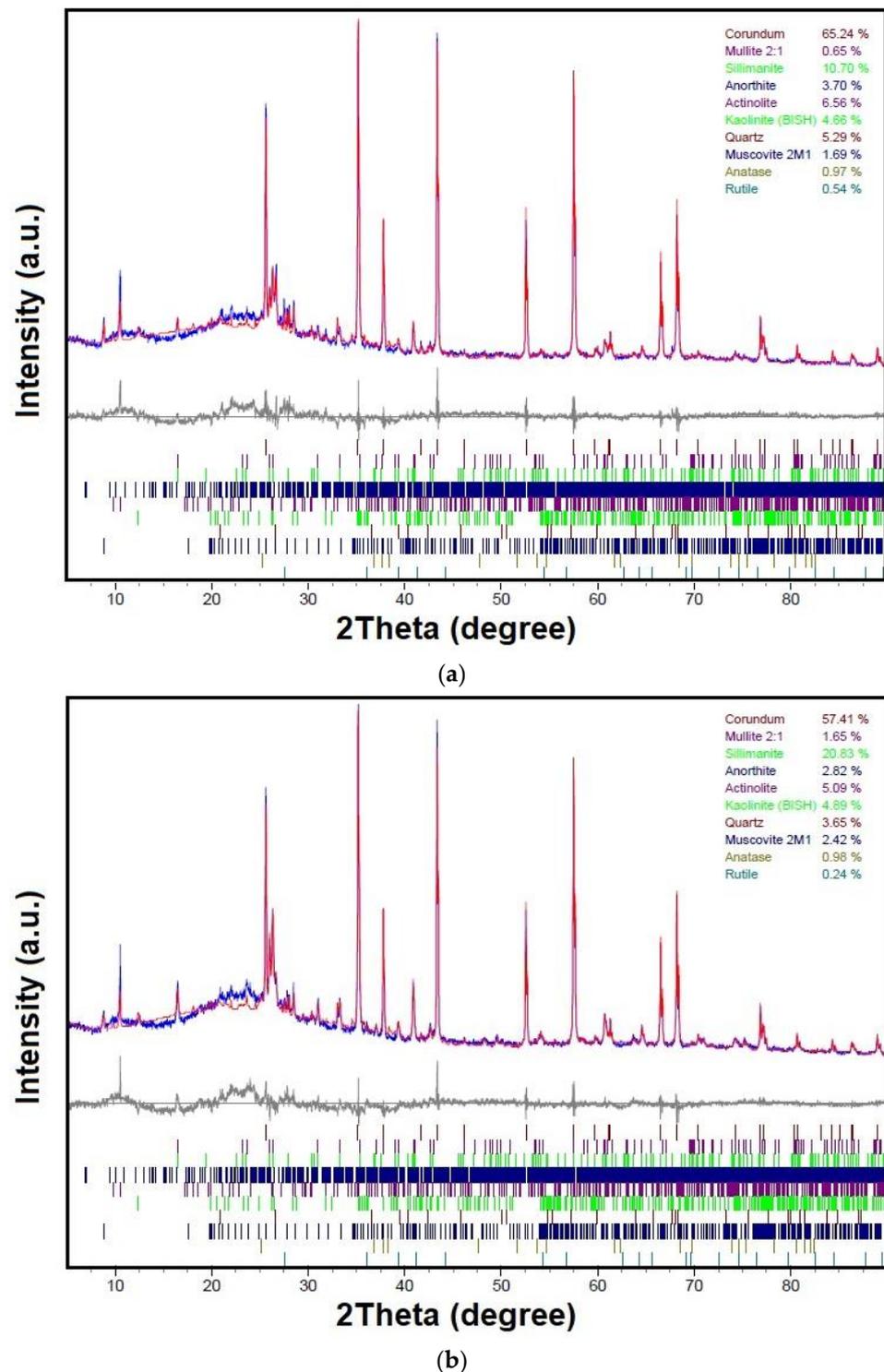


Figure 14. Rietveld refinement plots. (a) Rietveld refinement pattern of a normal porcelain insulator prototype; (b) Rietveld refinement pattern of a nano-TiO₂ coated porcelain insulator prototype.

4.3. Microstructural Analysis

Figures 15 and 16 depict the sintered microstructure that corresponds to the prototype porcelain insulator design based on international standards and the outdoor insulator composition used in industrial-scale railway electrification systems. Using the SEM and EDS methods in the back-scattered electron mode, and spectrum and quantity analysis

were measured by TESCAN, Model: MIRA3. We compared the porcelain insulators with traditional coatings in Figure 15 and nano-TiO₂ coatings in Figure 16.

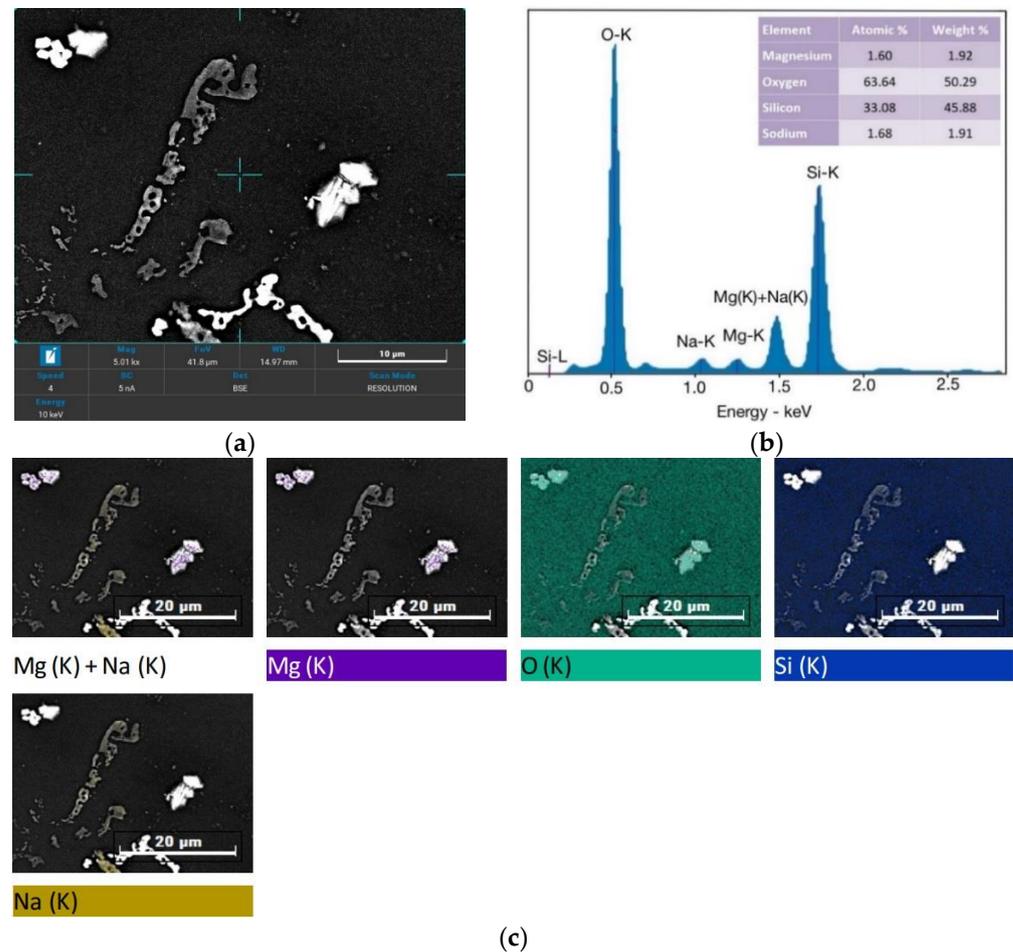


Figure 15. SEM and EDS of porcelain insulators with normal coatings. (a) SEM image of surface; (b) The EDS-spectrum and quantity analysis; (c) The element images.

In a glassy matrix with a consistent distribution of quasi-spherical pores, coarse quartz grains with variable shapes make up a typical siliceous porcelain microstructure. There are noticeable fissures all around the quartz grains. This occurrence is linked to the sample cooling stage after the prototype insulator has been burnt at 1300 °C for 60 h in a temperature-controlled kiln when the quartz grain volume falls by 2% because of the β - α quartz-phase transformation [7,31]. This shift may cause enough strain to cause both the glassy matrix and the quartz grains to crack because of the large difference in thermal expansion between the quartz grains, which have a coefficient of thermal expansion of $23 \times 10^{-6} \text{ C}^{-1}$, and the glassy phase that surrounds them, which has a coefficient of thermal expansion of $3 \times 10^{-6} \text{ C}^{-1}$ (at temperatures between 20 and 700 °C). The size of the quartz particles and the rate at which they cool cause a reduction in the porcelain's mechanical and dielectric strength [31].

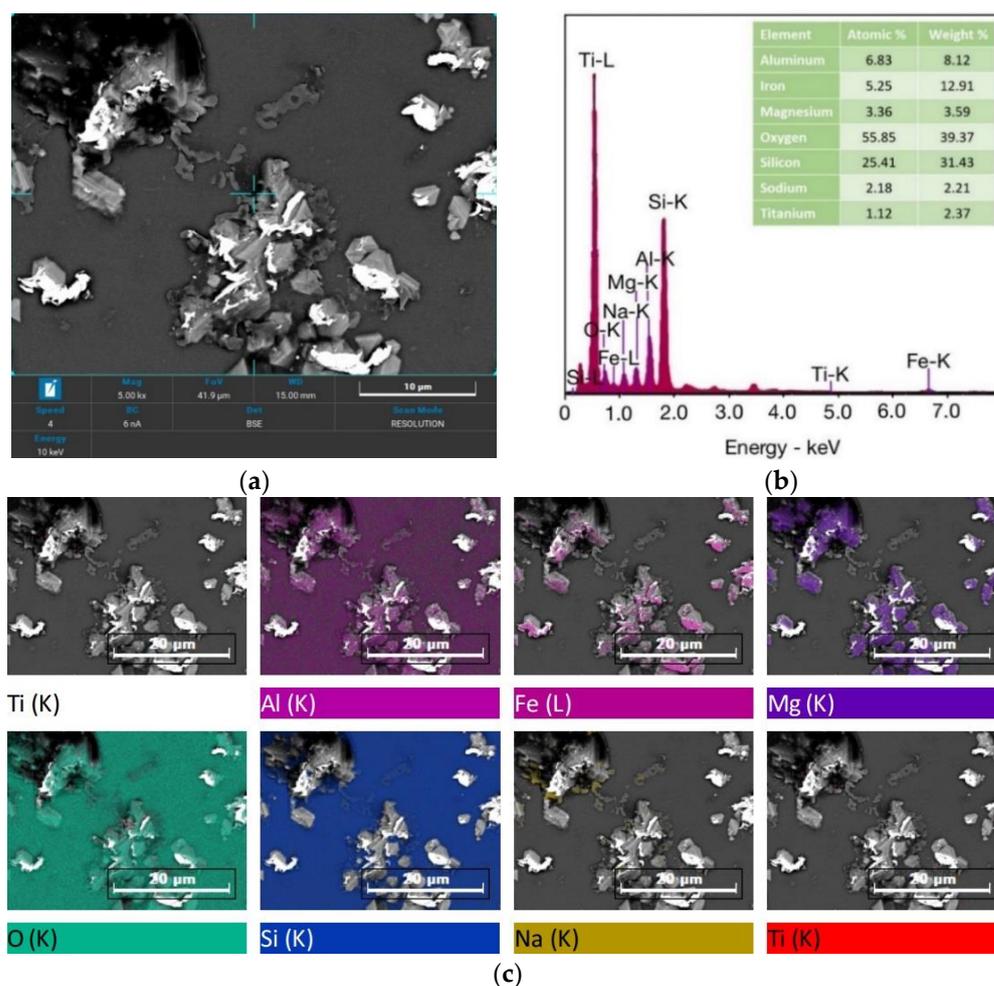


Figure 16. SEM and EDS of porcelain insulators with nano-TiO₂ coatings. (a) SEM image of surface; (b) The EDS-spectrum and quantity analysis; (c) The element images.

5. Conclusions

The test results concluded that when adding nano-TiO₂ at 0.1 wt.% of the amount of deionized water applications to outdoor insulators on railway electrification systems before processing coating and electrical testing, the performance of low-frequency flashover in dry and wet test conditions of the insulator coated with TiO₂ was higher than that coated with the normal coating. Furthermore, the test results of the lightning impulse critical-flashover voltage performance under normal test conditions showed that the insulator coated with nano-TiO₂ can cause a small change in the impulse breakdown voltage with both positive and negative polarity. Similarly, the positive polarity testing under contaminated test conditions, especially in the negative polarity of the impulse breakdown voltage, shows no improvement. However, it may be because the TiO₂ content is so low (only 0.1 wt.%) that it does not show a clear effect on the electrical properties of the coated insulators with improved coating as much as it should. In the future, we may add more TiO₂ to show higher electrical values. Finally, the results of this research showed that adding nano-TiO₂ to improve the electrical properties of the insulator will likely improve its performance in the future.

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