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# Green waste recycling of peanuts highly contaminated with aflatoxins in clay brick manufacturing

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## Abstract

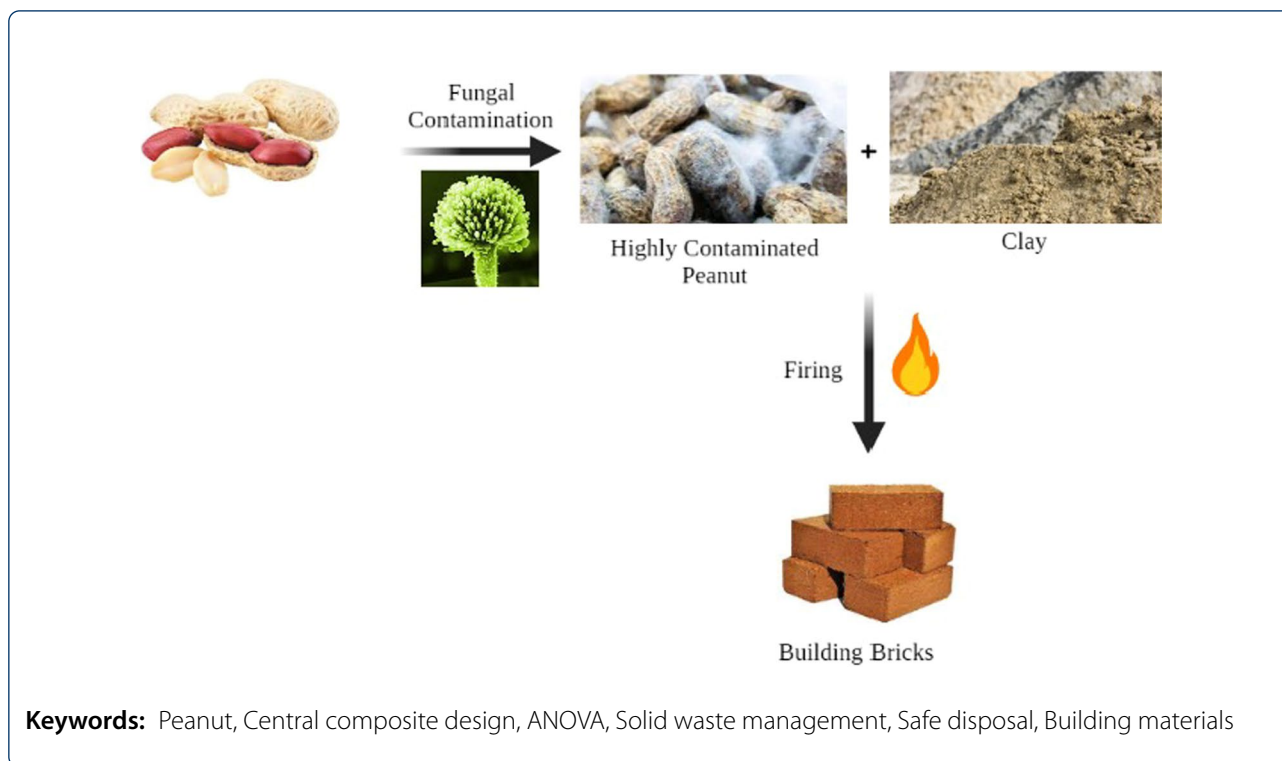
**Background:** The safe disposal of green waste is necessary for a clean environment, with the safe disposal of waste contaminated with aflatoxins being of particular importance. This work presents a novel route for utilizing contaminated grains in the production of clay bricks. In this work, the contaminated peanut (P) with aflatoxins (AFs) is divided into two categories: PA is the contaminated peanut kernel (without the outer shell), and PB is contaminated whole peanut grain (with the outer shell). Both of ground PA and PB were used to replace the clay in fired building bricks. The raw materials were characterized using X-ray fluorescence (XRF), X-ray diffraction (XRD), and thermo-gravimetric analyses (TGA). The effectiveness of this approach was evaluated by measuring the effect of the replacement percentage and firing temperature on the properties of the produced building bricks.

**Results:** The optimization of the process parameters was performed using central composite design as a tool in the response surface methodology. The ANOVA analysis of the predicted quadratic model elucidated significant models' terms and adequate precision that emphasizes the applicability of the model to navigate the design space. The results revealed that the optimum conditions are 6% clay replacement and 725 °C firing temperature, resulting in a compressive strength of 109.85 kg cm<sup>-2</sup> for clay bricks with PA and 126.33 kg cm<sup>-2</sup> for clay bricks with PB.

**Conclusions:** The objective of clay replacement by the contaminated peanut is not to improve the clay brick properties but to perform safe disposal of the accumulated contaminated peanut without deviating from the standards. The design of experiment using response surface method enables studying the effect of several factors on the clay brick properties simultaneously. Subsequently, this approach elucidates a sustainable route for accumulated contaminated green peanut waste disposal as the other alternatives have realized risks.

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## Highlights

- A new, green, and novel disposal of Aflatoxins in peanut was presented in the current study as additive in bricks production.
- The produced green bricks will withstand handling with negligible losses.
- This study paves the way for a more sustainable environment with zero-waste production.

## Background

Aflatoxins (AFs) are mycotoxins produced by toxigenic molds, such as *Aspergillus flavus* and *Aspergillus parasiticus*, the most toxic types of which are B1, B2, G1, and G2. AFs are highly genotoxic carcinogenic compounds, classified as a Group I carcinogen by the International Agency for Research on Cancer (IARC) (Saha and Wu 2019). AFs-producing fungi are predominantly found in hot and humid climate zones and their presence in food can be due to both preharvest and postharvest fungal contamination. The level and degree of contamination depend on the temperature, moisture content, soil type, and storage conditions (Schrenk et al. 2020). AF contamination is very common in many foods, including peanuts, nuts, spices, oilseeds, and products of these

foodstuffs (Mohammed et al. 2018; Al Ayoubi et al. 2021; Kortei et al. 2021). Various health risk assessment studies have been published recently regarding AFs worldwide; these were prompted due to health concerns and alerts for potential health effects on humans (Wang et al. 2018; Sebaei et al. 2020). Significantly, more rigid guidelines have been set up by the European Commission (EC) with a maximum permissible limit (MPL) of  $4 \mu\text{g kg}^{-1}$  for AFs in peanuts, dried organic products, and cereals for direct human consumption or use as food ingredients (EC, 2010). All shipments exceeding the MPL are rejected, and many of them represent a health risk when returning to exporting countries, especially developing countries.

Because of the extreme stability of AFs under different harsh conditions, including high heat and extreme pH (measurement of pH below about 2.5, and above about 10.5), it is very hard to eliminate AFs completely from food and fodder. Various techniques to minimize the risk of exposure to AFs have been reported. These approaches can be classified as a preharvest counteraction to the creation of AFs and postharvest AF reduction or detoxification. The use of good manufacturing practices, i.e., the utilization of fertilizers, pest control, antifungal crop varieties, harvesting at the opportune time, and keeping up low dampness and temperature during marketing, can reduce fungal growth and the production of AFs in grains. These methodologies are generally unsuccessful

in the elimination of AFs from food and fodder (Guo et al. 2021). This emphasizes the need for postharvest elimination of AF contamination. Traditional postharvest reduction techniques to detoxify AFs include physical and chemical treatments, such as irradiation, extrusion cooking, ozone treatment, reaction with organic acids, and ammoniation. Meanwhile, biocontrol by microorganisms has acquired prominence because of its friendliness to both the environment and humans (Peng et al. 2018).

Several attempts at waste utilization and recycling have been conducted in numerous countries worldwide to achieve reduced ecological impact or increased economic savings (Gupta et al. 2020). Some interesting research has combined different waste products into clay bricks as a substitute, additive, or support; this research has frequently shown that the concept is of positive value both economically and environmentally. The use of organic waste additives has also been recommended by several authors because of the exothermic character of their oxidation, which represents a key factor in decreasing the fuel required for firing.

Such works have used many waste products, including waste tea (Demir, 2006), cigarette butts (Abdul-Kadir and Mohajerani 2008; 2011), dried mango leaves (Folaranmi, 2009), pineapple leaves and oil palm fruit bunch (Chan 2011), sugarcane bagasse ash (Faria et al. 2012), plastic waste (Mobili et al. 2018), travertine sludge (Shishегaran et al. 2021), wastewater treatment plant sludge (Limami et al. 2021), industrial waste (Balasubramaniam et al. 2021), waste products from concrete (Mahmoodi et al. 2021), industrial acetylene biomass ash (Eliche-Quesada et al. 2021), potato peel powder and sour orange leaf (Ghorbani et al. 2021), coal dust (Vasić et al. 2021), date palm waste (Khouidja et al. 2021), and peanut shell ash (Diedhiou et al. 2019).

The utilization of whole contaminated grains in the production of fired clay bricks has not been reported previously. In this work, the evaluation of this novel approach has been undertaken by investigating the properties of the fired bricks and the effect of the replacement percentage and the firing temperature on the product characteristics. The optimization of these parameters was performed using central composite design (CCD) as a tool for use in the response surface methodology (RSM) to minimize the number of experiments (El-Mekkawi et al. 2020).

The present work investigates the possibility of utilizing peanuts contaminated with AFs in the manufacture of clay building bricks. Such a method would constitute a step toward an integrated sustainable solution for diminishing the harmful effect of contaminated grains,

providing an ecofriendly product, and preserving clay as a natural resource. In this study, peanuts contaminated with AFs are divided into two groups. The first group (PA) is the contaminated peanut kernel (without the outer shell), and the second (PB) is the contaminated whole peanut grain (with the outer shell). Ground PA and PB are added to the clay in certain proportions and fired at scheduled temperatures to examine their effect on the clay brick characteristics.

## Methods

### Raw materials

#### *Peanuts*

Ten kilograms of contaminated peanut samples without their shell (peanut kernel) and with their shell was used in this experiment. The peanut stock was collected from local retail sources. The peanuts were analyzed for AFs in the Central Laboratory of Pesticides and Heavy Metals in Food, which showed high contamination levels with an AF concentration exceeding 100 µg/kg. The two samples were crushed and blended for homogenization using a high-speed blender; the samples after preparation had a particle size smaller than 2 mm. They were then tested for the main chemical composition and ash content using the standard test method AOAC 923.03 (2016), for moisture content following the test procedure AOAC 925.09 (2016), and for fat content according to the standard AOAC 2003.05 (2016). A chemical analysis of the two peanut wastes was undertaken using an X-ray fluorescence (XRF) diffractometer (Philips PW 1730).

#### *Clay brick basic mix*

The clay material used in the manufacture of the clay bricks was obtained from Arab Abu-Saed, Helwan, Cairo, Egypt. The clay material used was subjected to the following analyses: The chemical composition was assessed using an XRF diffractometer (Philips PW1730), the mineralogical composition was obtained using a computerized X-ray diffractometer (Bruker D8), and a thermal analysis was conducted using a combined DTA and TGA apparatus (Netzsch STA 409 C/CD) at a heating rate of 10 °C min<sup>-1</sup> in nitrogen. The clay material was screened according to ASTM D 422 (2007) to estimate its particle size distribution using a standard set of sieves in agreement with ASTM E 11 (2021). The screen apertures varied from 4.75 mm (4 mesh) to 45 µm (325 mesh). Finally, the free silica content was determined via the standard method of Kumari and Mohan (2021), and the powder density was determined using the liquid pycnometer method (ASTM D 854/2014 2021; ISO 1183-1/2019).

**Table 1** Matrix for the CCD

Factor	Level				
	Low	Central	High		
	(-1)	(0)	(+1)	(-α)	(+α)
X <sub>1</sub> : Clay replacement, %	6	20.50	35	0	41
X <sub>2</sub> : Firing temperature, °C	725	800	825	693.93	906

**Design of experiments**

The simultaneous optimization of the degree of clay replacement (whether using sample PA or PB) and the firing temperature is necessary to investigate any possible interaction between these two parameters and to optimize the compressive strength of the fired brick sample. With this aim, 13 duplicated experiments were performed on the basis of a range of replacement percentages (6%–35%) and a range of firing temperatures (725–875°C) using central composite design (CCD) in the response surface methodology (RSM) by the Design Expert 6.0.8 software during a trial period. The specified ranges, as shown in Table 1, were selected on the basis of preliminary experiments. The parameters used in this study are given in Table 1. Each parameter was analyzed statistically at five levels; these levels were coded as -α, -1, 0, +1, +α, respectively. The total number of experiments undertaken was 13, with four factorial points (2<sup>2</sup>), four axial points (2 × 2), and five replicate points in accordance with the CCD.

The mathematical relationship between the parameters and the response can be approximated using the following second-order polynomial model (El-Mekkawi et al. 2020):

$$Y = a_0 + a_1X_1 + a_2X_2 + a_{11}X_1^2 + a_{22}X_2^2 + a_{12}X_1X_2 \tag{1}$$

where *Y* is the predicted response; *a*<sub>0</sub> is the intercept regression constant; *a*<sub>1</sub>, *a*<sub>2</sub>, ..., *a*<sub>12</sub> are regression coefficients; and *X*<sub>1</sub> and *X*<sub>2</sub> are the independent variables investigated. Analysis of variance (ANOVA) was performed to judge the significance and goodness of fitting using the developed model.

**Preparation of the brick samples**

The set of 13 triplicated cubic samples were molded with dimensions of 50 × 50 × 50 mm<sup>3</sup> by dry pressing using a laboratory hydraulic press under uniaxial pressure of 40 kg cm<sup>-2</sup> with 15% water as a binder. The samples were dried in three phases using a laboratory dryer. The first phase was at 50 °C for 24 h, followed by drying at 80 °C for 3 h, and finally drying at 110 °C for 3 h. The linear shrinkage and green compressive strength of the unfired brick samples were determined according to the ASTM

standards (ASTM C 326/2018 2021; ASTM C 67/2020 2021). The samples were fired at the five temperatures stated in Table 1, using a Prothem electrical furnace model PLF 14,015 with a heating rate of 10 °C/min and a total firing time of 2 h.

**Determination of the properties of the fired bricks**

The physical and mechanical properties of the fired clay brick samples were tested according to the ASTM standards. The firing shrinkage was measured according to ASTM C 326 (2021), whereas the cold and boiling water absorption as well as the saturation coefficient were calculated following ASTM C 67 (2021). The bulk density and apparent porosity were determined according to ASTM C 20 (2015). Finally, the compressive strength was determined following ASTM C 67 (2021).

**Results**

**Raw materials characterization**

**Peanuts**

Table 2 addresses the compositions of moisture, ash, and oil in the two contaminated peanut samples, PA and PB. The bricks using the additive PA showed a lower ash and a higher oil content than did the bricks using the additive PB.

The chemical composition of the two samples, as determined by the XRF analysis, is presented in Table 3. Generally, the higher silica content in PB is due to the peanut shell composition, which is present in sample PB but not in PA, and the elevated loss on ignition of both PA and PB is due to the high content of organic compounds in both samples.

**Clay brick basic mix**

The chemical composition of the clay material used was obtained via XRF analysis and is given in Table 4. This table shows that the main components present are SiO<sub>2</sub>, Al<sub>2</sub>O<sub>3</sub>, and Fe oxides, which account for approximately 86% of the composition. Minor oxides also present include alkaline earth oxides (CaO and MgO) and alkali oxides (Na<sub>2</sub>O and K<sub>2</sub>O). The SO<sub>3</sub> appearing in the analysis is related to the presence of small quantities of gypsum (CaSO<sub>4</sub>·2H<sub>2</sub>O), anhydrite (CaSO<sub>4</sub>), or MgCO<sub>3</sub>.

**Table 2** Composition of the two peanut additive samples: ash, oil, and moisture contents (mean values of triplicated runs)

	Ash content (%)	Oil content (%)	Moisture content (%)
PA	2.6	47.6	5.3
PB	3.0	34.0	6.2

**Table 3** Chemical analysis of the two contaminated peanut samples

Main constituent	PA (W%)	PB (W%)
SiO <sub>2</sub>	0.56	0.97
Al <sub>2</sub> O <sub>3</sub>	0.18	0.32
Fe <sub>2</sub> O <sub>3</sub> tot	0.01	0.01
MgO	0.16	0.21
CaO	0.13	0.14
Na <sub>2</sub> O	0.15	0.12
K <sub>2</sub> O	0.27	0.25
P <sub>2</sub> O <sub>5</sub>	0.34	0.16
SO <sub>3</sub>	0.28	0.17
Cl	0.1	0.05
ZnO	0.002	0.003
SrO	0.001	–
HfO <sub>2</sub>	0.007	–
LOI	97.78	97.58
Total	99.97	99.983

**Table 4** Chemical analysis of the clay material

Main constituent	(wt%)
SiO <sub>2</sub>	58.67
Al <sub>2</sub> O <sub>3</sub>	14.69
Fe oxides	6.71
CaO	3.46
MgO	1.38
Na <sub>2</sub> O	1.62
K <sub>2</sub> O	1.07
TiO <sub>2</sub>	0.88
Cl <sup>-</sup>	1.14
SO <sub>3</sub>	3.40
Others	0.04
LOI	7.02
Total	100.08

The X-ray diffraction (XRD) analysis of the clay material indicates that the clay raw material is mainly composed of kaolinite (Al<sub>2</sub>O<sub>3</sub>·2SiO<sub>2</sub>·2H<sub>2</sub>O), quartz (SiO<sub>2</sub>), montmorillonite (Na<sub>x</sub>(Al,Mg)<sub>2</sub>Si<sub>4</sub>O<sub>10</sub>(OH)<sub>2</sub>·zH<sub>2</sub>O), and hematite (Fe<sub>2</sub>O<sub>3</sub>), as shown in Fig. 1. Other probable compounds, including calcium carbonate, magnesium carbonate, sulfates, and chlorides, were not detected because of their low percentages. These results are incongruent with those obtained via XRF analysis (Rahaman, 2003; Földvári, 2011).

The DTA trace of the clay material used in this work is shown in Fig. 2. It shows three principle endothermic peaks besides the first peak at approximately 100 °C,

which is associated with the elimination of moisture: The first peak at about 270 °C is hypothesized to correspond to the decomposition of gypsum to anhydrite and magnesium carbonate to magnesium oxide (Rahaman, 2003; Földvári, 2011), the second peak starting at approximately 400 °C and ending at approximately 600 °C corresponds to the dehydroxylation of kaolinite, and finally the minor peak at approximately 870 °C corresponds to the calcination of calcium carbonate. All these peaks are accompanied by weight losses, which can be observed in the TGA–DTG traces shown in Fig. 3.

The calculated weight loss excluding that due to physical moisture loss was approximately 4.4%, implying the presence of a large quantity of free silica. If all the available alumina was assumed to be present in kaolinite, then the combined silica would represent approximately 17.3% of the total silica present, and the available kaolinite would make up 37.1%. This leaves approximately 41.4% as free quartz. Because the mass of the dry clay sample used in the TGA was 15.08 mg, then the calculated amount of free silica was 6.24 mg and the mass of kaolinite was 5.59 mg. The loss in weight for dehydroxylation was found to be 0.53 mg, representing a loss of approximately 9.5%, a typical figure for weight loss of kaolin minerals. The fact that the actual LOI is approximately 7% is hypothesized to be due to kaolinite being assumed to have the model structure, which neglects the possibility of impurities, such as iron oxides.

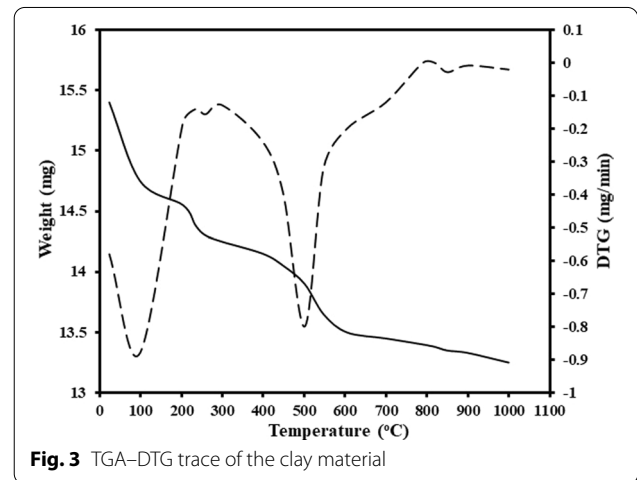
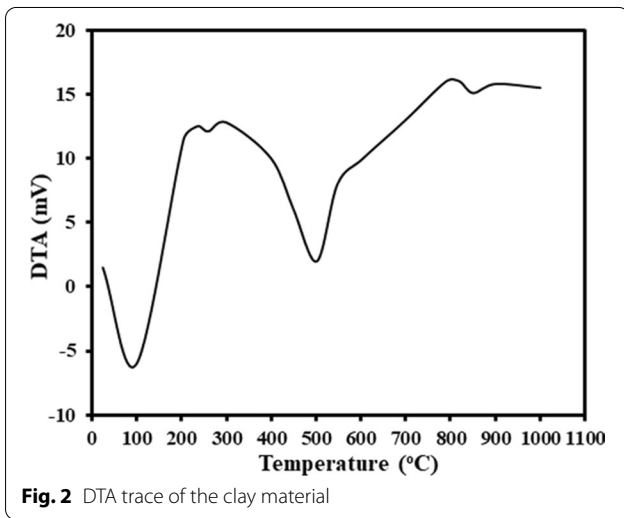
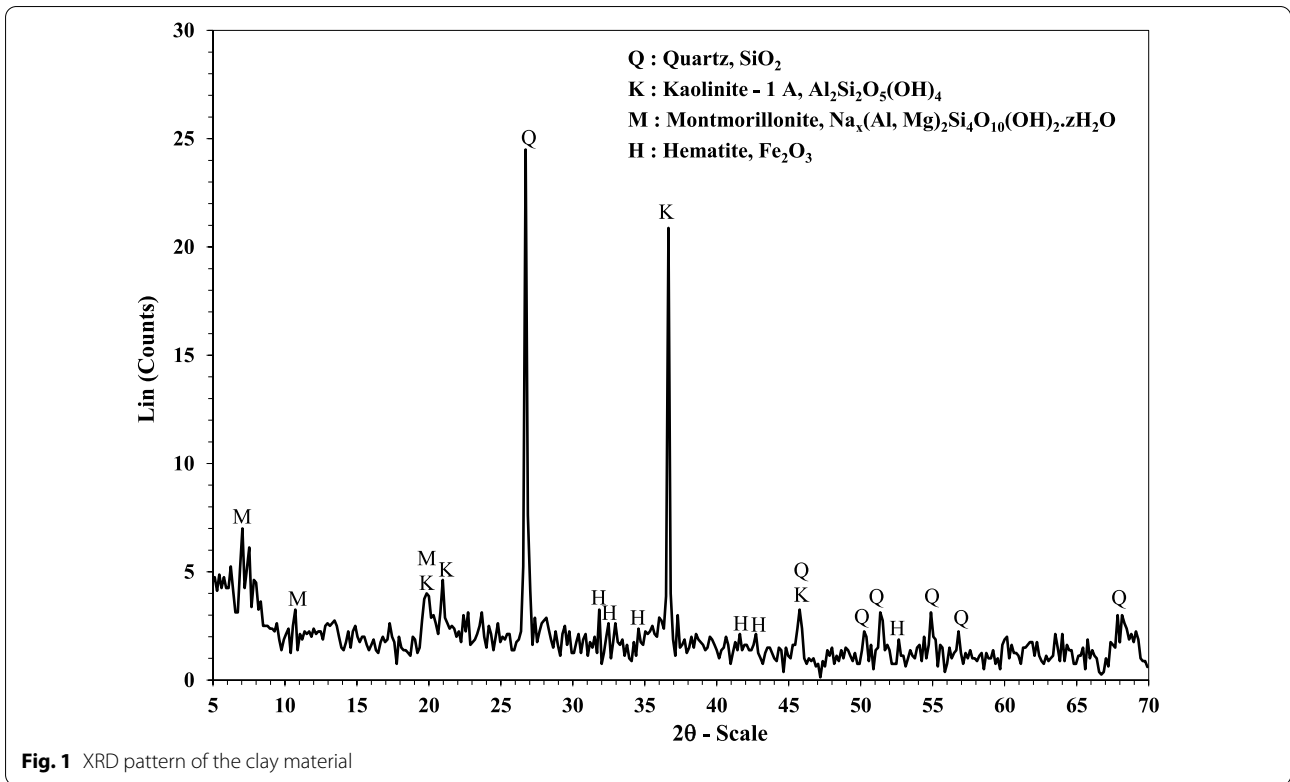
The particle size distribution of the clay material used is shown in Fig. 4. It reveals a median particle size, *D*<sub>50</sub>, of 0.31 mm. The bimodal nature of the curve is due to the presence of free silica, which is harder to grind than kaolin. The first peak is found at a particle size of approximately 0.16 mm, and the second peak, corresponding to quartz, can be observed corresponding to a particle size of 1 mm.

Following Mc Cabe et al. (2005), the mean particle size of a powder having the same specific surface area of the material can be calculated according to:

$$D_{vs} = \frac{1}{\sum_{i=0}^n \frac{x_i}{D_{avi}}} \tag{2}$$

where *n* represents the number of sieves, *x*<sub>*i*</sub> represents the mass fraction of solid retained between sieve *i*–1 and *i*, and *D*<sub>avi</sub> represents the average screen opening between sieve *i*–1 and *i*. Using this method, the mean particle size was found to be 0.183 mm.

Finally, the free silica content of the clay material used in this study was found to be 41%. This implies that it was not necessary to add any sand to the clay in the brick mix. The powder density was found to be 2.58 g cm<sup>-3</sup>.



**Characterization of green bodies**

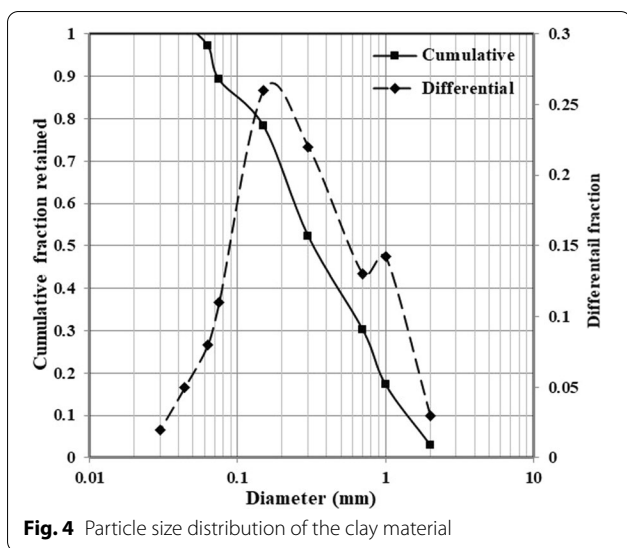
**Linear drying shrinkage**

The results shown in Fig. 5 indicate a significant increase in the linear drying shrinkage percentage for both the clay-PA and clay-PB compared with the control clay specimen without the addition of waste products. This increase is a consequence of the presence of oil, which facilitates the motion of the constituent particles during

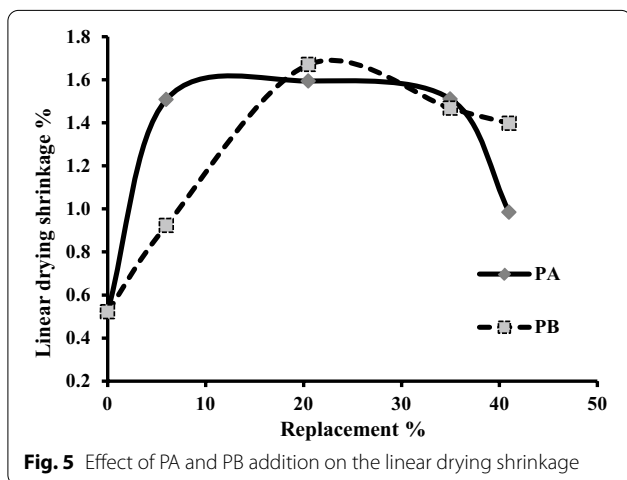
shrinkage. The stability against shrinking is enhanced by increasing the ash content and inhibited by increasing the oil content in the bricks. The general trend of decreasing shrinkage on increased replacement is due to the non-plastic nature of peanuts, the effect of which is greater than the effect of the oil.

**Green compressive strength**

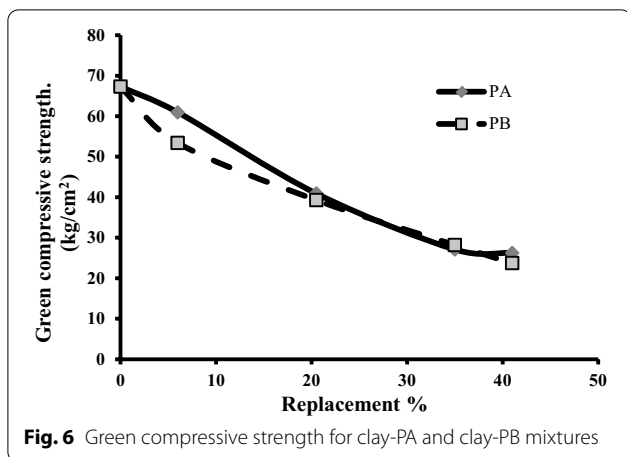
Green compressive strength does not have a standard limitation; however, it indicates the capacity of the



**Fig. 4** Particle size distribution of the clay material



**Fig. 5** Effect of PA and PB addition on the linear drying shrinkage



**Fig. 6** Green compressive strength for clay-PA and clay-PB mixtures

**Table 5** Compressive strength of clay-PA and clay-PB fired bricks

Run	$X_1$	$X_2$	Compressive strength of clay-PA bricks (Model I)	Compressive strength of clay-PB bricks (Model II)
1	6.00	725.00	109.85	126.33
2	35.00	725.00	5.32	6.33
3	6.00	875.00	66.27	70.09
4	35.00	875.00	5.10	6.31
5	0	800.00	339.30	339.30
6	41.01	800.00	5.09	5.74
7	20.50	693.93	13.89	14.71
8	20.50	906.07	10.35	9.00
9	20.50	800.00	11.80	11.97
10	20.50	800.00	12.90	11.70
11	20.50	800.00	12.00	11.50
12	20.50	800.00	12.75	12.90
13	20.50	800.00	12.20	12.20

specimen to be formed and handled without failure. Figure 6 shows a linear decrease in the green compressive strength as the replacement percentage of the two mixtures increases. The non-plastic nature of both waste products will induce a lack of cohesion between the constituent particles and hence decrease the green strength. This severe increase is due to the significant oil content,  $47.6\% \pm 1.14\%$  of PA and  $34.0\% \pm 0.81\%$  of PB. The minimum strength observed in both cases exceeds  $20 \text{ kg cm}^{-2}$ . This figure is high enough to suggest that the green bricks will withstand handling with negligible losses.

**Discussion**

The compressive strength was determined experimentally to determine the fired bodies characterization. The clay-PA and clay-PB fired bricks at the chosen values of the parameters  $X_1$  and  $X_2$ , as illustrated in Table 5. The experiments were analyzed using the Design-Expert software. Two predicted models were obtained that exhibit significant curvature for both the clay-PA (model I) and clay-PB (model II) fired bricks through the design points. This finding has required supplementary axial points to be added to fit the models.

**Regression model for clay-PA fired bricks (model I)**

A logarithmic quadratic model was suggested by the software, as shown in Table 6. The regression coefficient  $R^2$  and the adjusted  $R^2$  are close to 1; the predicted  $R^2$  is in reasonable agreement with the adjusted  $R^2$ . Both the linear and quadratic models are valid when the probability  $F$ -value ( $p$ ) is less than 0.05. However, the error function

**Table 6** Suggested model summary statistics

Source	SD <sup>a</sup>	R <sup>2</sup>	R <sup>2</sup> Adj <sup>b</sup>	R <sup>2</sup> Pred <sup>c</sup>	PRESS <sup>d</sup>	SS Prob>F <sup>e</sup>	Lack of fit Prob>F <sup>f</sup>	
Linear	0.51	0.8649	0.8379	0.7345	5.14	<0.0001	<0.0001	
2F1	0.53	0.8677	0.8236	0.7142	5.53	0.6746	<0.0001	
Quadratic	0.065	0.9985	0.9974	0.9908	0.18	<0.0001	0.0698	Suggested
Cubic	0.050	0.9994	0.9984	0.9778	0.43	0.1149	0.1034	Aliased

<sup>a</sup> Standard deviation

<sup>b</sup> Adjusted residual squared

<sup>c</sup> Predicted residual squared

<sup>d</sup> Predicted residual error sum of squares

<sup>e</sup> Sequential model sum of the squares probability function

<sup>f</sup> Lack of fit probability function

**Table 7** ANOVA results for the CCD

Source	Sum of squares	df	Mean square	F-value	Prob>F	
Model	19.31	4	4.83	1099.53	<0.0001	s
X <sub>1</sub>	16.62	1	16.62	3784.43	<0.0001	s
X <sub>2</sub>	0.12	1	0.12	26.43	0.0009	s
X <sub>1</sub> <sup>2</sup>	2.52	1	2.52	575.04	<0.0001	s
X <sub>1</sub> .X <sub>2</sub>	0.054	1	0.054	12.20	0.0082	s
Residual	0.035	8	4.391E-003			
Lack of fit	0.029	4	7.296E-003	4.91	0.0761	e
Pure error	5.941E-003	4	1.485E-003			

s Significant

e Nonsignificant

is significant for all the models except for the quadratic model. All the terms of the quadratic model are significant according to the ANOVA undertaken, which is summarized in Table 7. The model terms are two linear factors (X<sub>1</sub>, X<sub>2</sub>), a quadratic factor X<sub>1</sub><sup>2</sup>, and the product (X<sub>1</sub>.X<sub>2</sub>). The coefficient of variance (CV) is 2.32, which suggests the relationships between the model terms could be predicted within the selected range.

The regression equation of the actual variables is

$$Y = 8.75313 - 0.30099X_1 - 0.00387X_2 + 0.00284X_1^2 + 0.0001064X_1.X_2 \tag{3}$$

where Y = ln σ, σ is the compressive strength and all other variables have been previously defined.

The notable curvature of the model output displayed in Fig. 7a demonstrates the necessity to perform runs 5–8 using the axial points to fit the model. There is no interaction between the parameters of the model in the design space for replacement levels lower than 35%, as shown in Fig. 7b. Plotting the residuals against the normal percent probability (Fig. 7c) illustrates a normal distribution through a linear relationship. The response actual

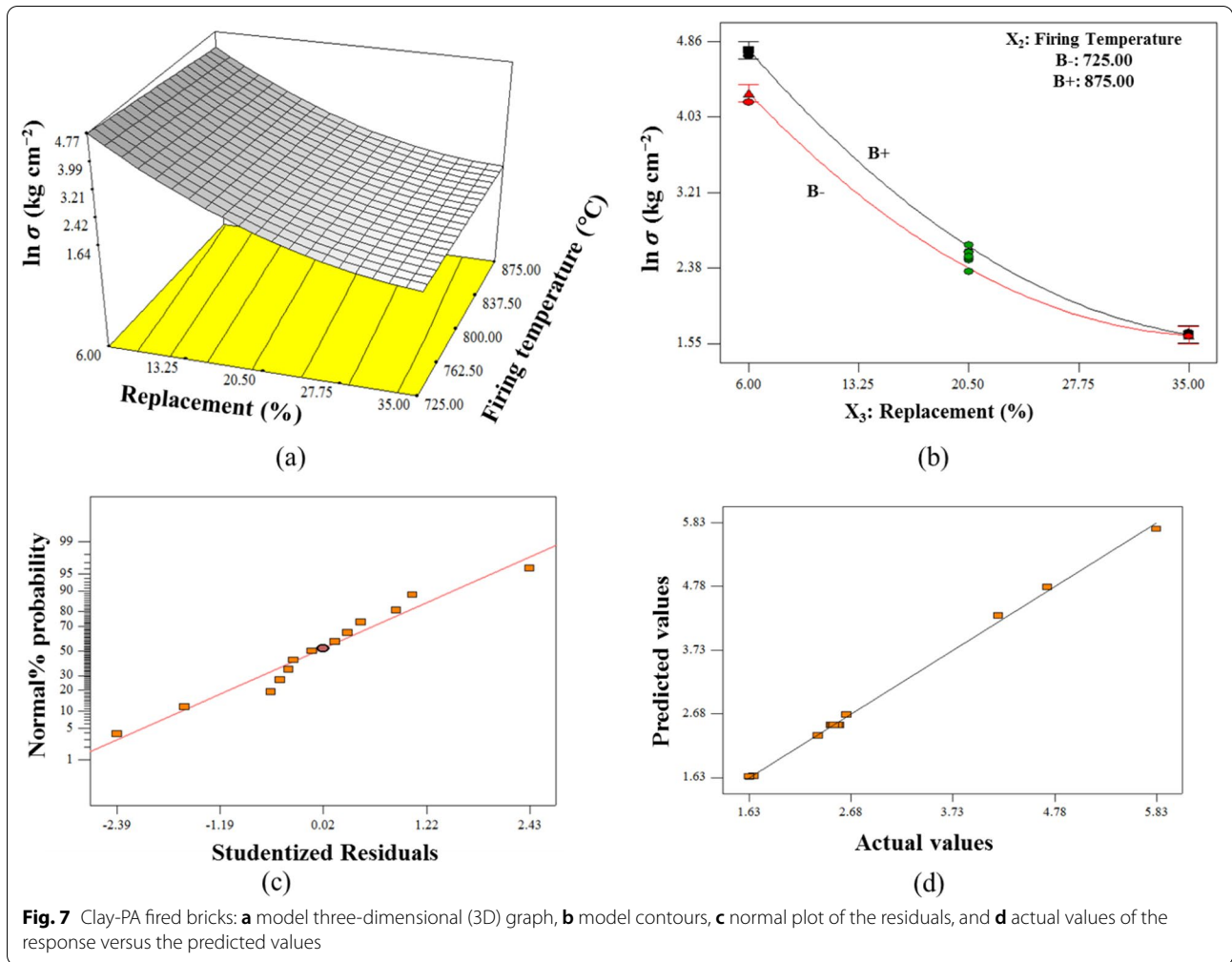
values and the response predicted values are congruent, as shown in Fig. 7d.

**Optimization of model I parameters**

The main objective of this model is to estimate the maximum compressive strength. This target is achieved at a 6% clay replacement and a 725 °C firing temperature. The predicted compressive strength under these conditions was found to be 117.57 kg cm<sup>-2</sup>, whereas the experimental value is 109.85 kg cm<sup>-2</sup>, meeting ASTM C 62 (2021), which requires a minimum compressive strength of 89.74 kg cm<sup>-2</sup>. This value is located within the 95% confidence interval (CI) and 95% prediction interval (PI), as illustrated in Table 8.

Other vitrification properties, such as loss on ignition (LOI), total linear shrinkage (TLS), cold water absorption (CWA), boiling water absorption (BWA), saturation coefficient (SC), and bulk density (BD), were determined to investigate the behavior of fired bricks with PA replacement. The regression equations that describe the relationship between the parameters and their responses are as follows:





**Table 8** Statistical analysis of the response

	Prediction	SE <sup>a</sup> mean	95% CI low	95% CI high	SE <sup>b</sup> pred	95% PI low	95% PI high
ln(Compressive strength)	4.767	0.051	4.65	4.88	0.084	4.57	4.96

<sup>a</sup> Standard error of the mean

<sup>b</sup> Standard error of the prediction

$$\begin{aligned}
 \text{LOI (\%)} = & -4.26976 + 0.60043X_1 + 0.016408X_2 \\
 & + 0.00738689X_1^2
 \end{aligned}
 \tag{4}$$

$$\text{TLS (\%)} = -6.06335 + 0.16034X_1 + 0.008125X_2
 \tag{5}$$

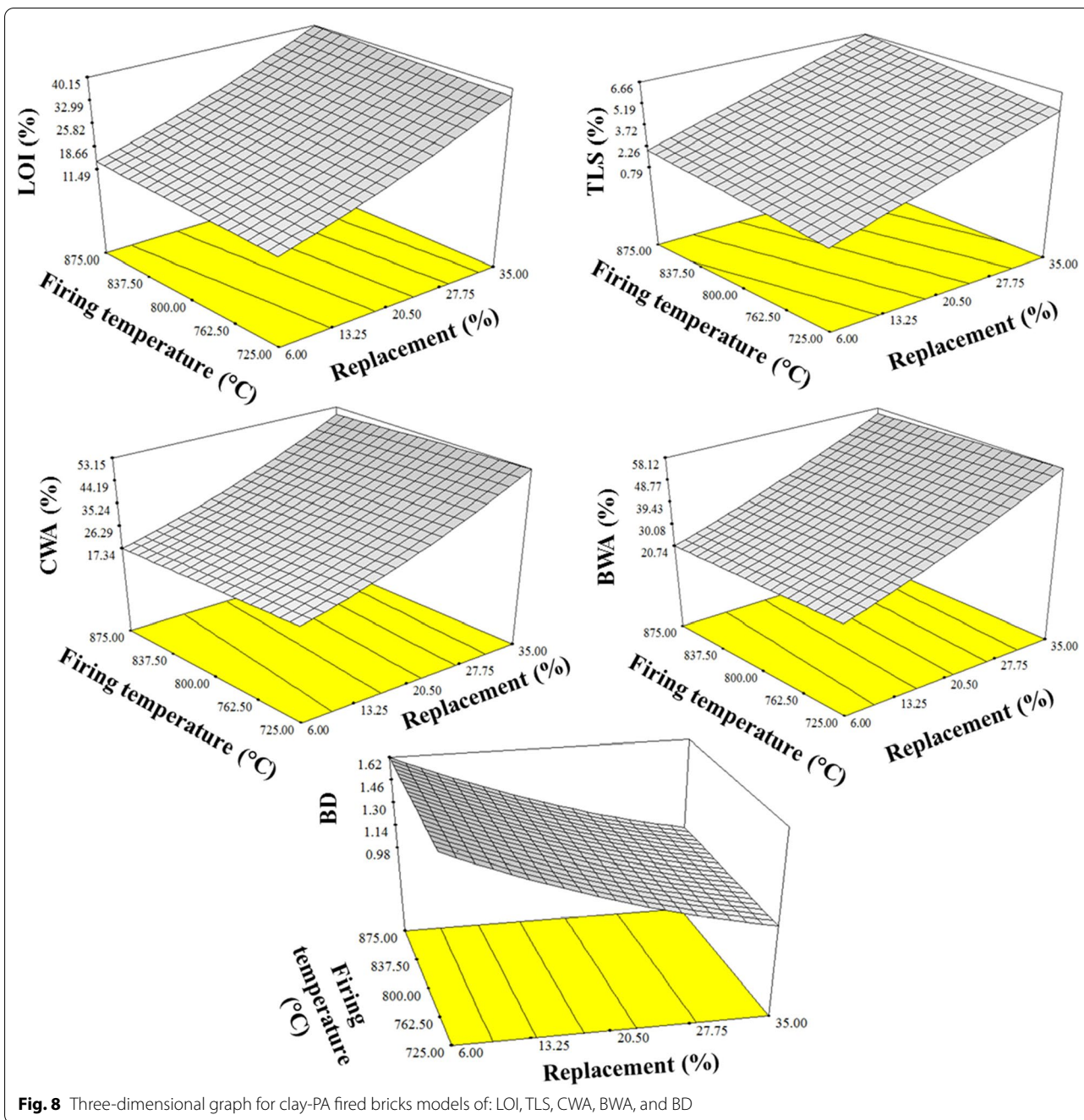
$$\begin{aligned}
 \text{CWA (\%)} = & 30.13513 + 0.58901X_1 \\
 & - 0.019213X_2 + 0.013327X_1^2
 \end{aligned}
 \tag{6}$$

$$\begin{aligned}
 \text{BWA (\%)} = & 28.40433 + 0.88266X_1 \\
 & - 0.015145X_2 + 0.00800103X_1^2
 \end{aligned}
 \tag{7}$$

$$\text{BD (g cm}^{-3}\text{)} = 1.81104 - 0.033724X_1 + 0.0002867X_1^2
 \tag{8}$$

It is noteworthy that, as illustrated in Fig. 8, according to the model, the LOI is a quadratic model that depends predominantly on the replacement percentage. However, the lowest LOI was required to maintain high compressive strength and no black core formation. This is because the black core was present for replacements > 10%. The minimum LOI was achieved at 6% clay replacement and 725 °C.

The TLS showed a linear dependence on both parameters, as can be noted from the regression equations,



although the level of clay replacement showed a stronger effect. The contour of the TLS model shows that the minimum response is located at the minimum values of both parameters. Using 6% clay replacement and firing at 725 °C resulted in a high BD and a suitable SC of 0.89.

**Regression model for clay-PB fired bricks (model II)**

As explained in model I, a quadratic model with a natural log was suggested by the software for model II (see

Table 9). The model terms of the quadratic model are significant, as shown in Table 10. The relationships between the model terms could be predicted within the selected range since the C.V. is 2.39.

The regression equation of the actual variables was found to be:

$$Y = 9.72112 - 0.33450X_1 - 0.00490306X_2 + 0.00318404X_1^2 + 0.000134568X_1.X_2 \tag{9}$$

**Table 9** Suggested model summary statistics (model II)

Source	SD <sup>a</sup>	R <sup>2</sup>	R <sup>2</sup> Adj <sup>b</sup>	R <sup>2</sup> Pred <sup>c</sup>	PRESS <sup>d</sup>	SS Prob>F <sup>e</sup>	Lack of fit Prob>F <sup>f</sup>	
Linear	0.57	0.8275	0.7930	0.6650	6.40	0.0002	< 0.0001	
2F1	0.60	0.8319	0.7759	0.6115	7.42	0.6358	< 0.0001	
Quadratic	0.074	0.9980	0.9966	0.9880	0.23	< 0.0001	0.0735	Suggested
Cubic	0.064	0.9989	0.9974	0.9578	0.81	0.2030	0.0663	Aliased

<sup>a</sup> Standard deviation

<sup>b</sup> Adjusted residual squared

<sup>c</sup> Predicted residual squared

<sup>d</sup> Predicted residual error sum of squares

<sup>e</sup> Sequential model sum of the squares probability function

<sup>f</sup> Lack of fit probability function

**Table 10** ANOVA results for the CCD (Model II)

Source	Sum of squares	df	Mean square	F-value	Prob>F	
Model	19.06	4	4.77	990.22	< 0.0001	S
X <sub>1</sub>	15.60	1	15.60	3241.10	< 0.0001	S
X <sub>2</sub>	0.21	1	0.21	43.00	0.0002	S
X <sub>1</sub> <sup>2</sup>	3.17	1	3.17	658.98	< 0.0001	s
X <sub>1</sub> .X <sub>2</sub>	0.086	1	0.086	17.80	0.0029	s
Residual	0.039	8	4.813E-003			
Lack of fit	0.031	4	7.651E-003	3.87	0.1090	e
Pure error	7.899E-003	4	1.975E-003			

s Significant

e Nonsignificant

A notable curvature of the model output is presented in Fig. 9a. There is no interaction between the parameters of the model in the design space for replacements lower than 35%, as shown in Fig. 9b. The normal residual graph is linear, as illustrated in Fig. 9c, and the response actual values and the response predicted values are congruent and form a linear relationship, as shown in Fig. 9d.

**Optimization of model II parameters**

The principal objective of this model is to obtain the maximum compressive strength. This target is achieved at a 6% clay replacement and a 725 °C firing temperature. The predicted compressive strength under these conditions is 128.895 kg cm<sup>-2</sup>, whereas the experimental value is 126.33 kg cm<sup>-2</sup>, thus exceeding ASTM C 62 (2021), which states that the minimum compressive strength is 89.74 kg cm<sup>-2</sup>. This value is located within the 95% CI and 95% PI, as illustrated in Table 11. The other investigated characteristics were LOI, TLS, CWA, BWA, SC, and BD. The regression equations of the model describe the relationship between the parameters and their responses is:

$$\text{LOI (\%)} = - 5.80453 + 0.67640X_1 + 0.017703X_2 + 0.00360192X_1^2 \tag{10}$$

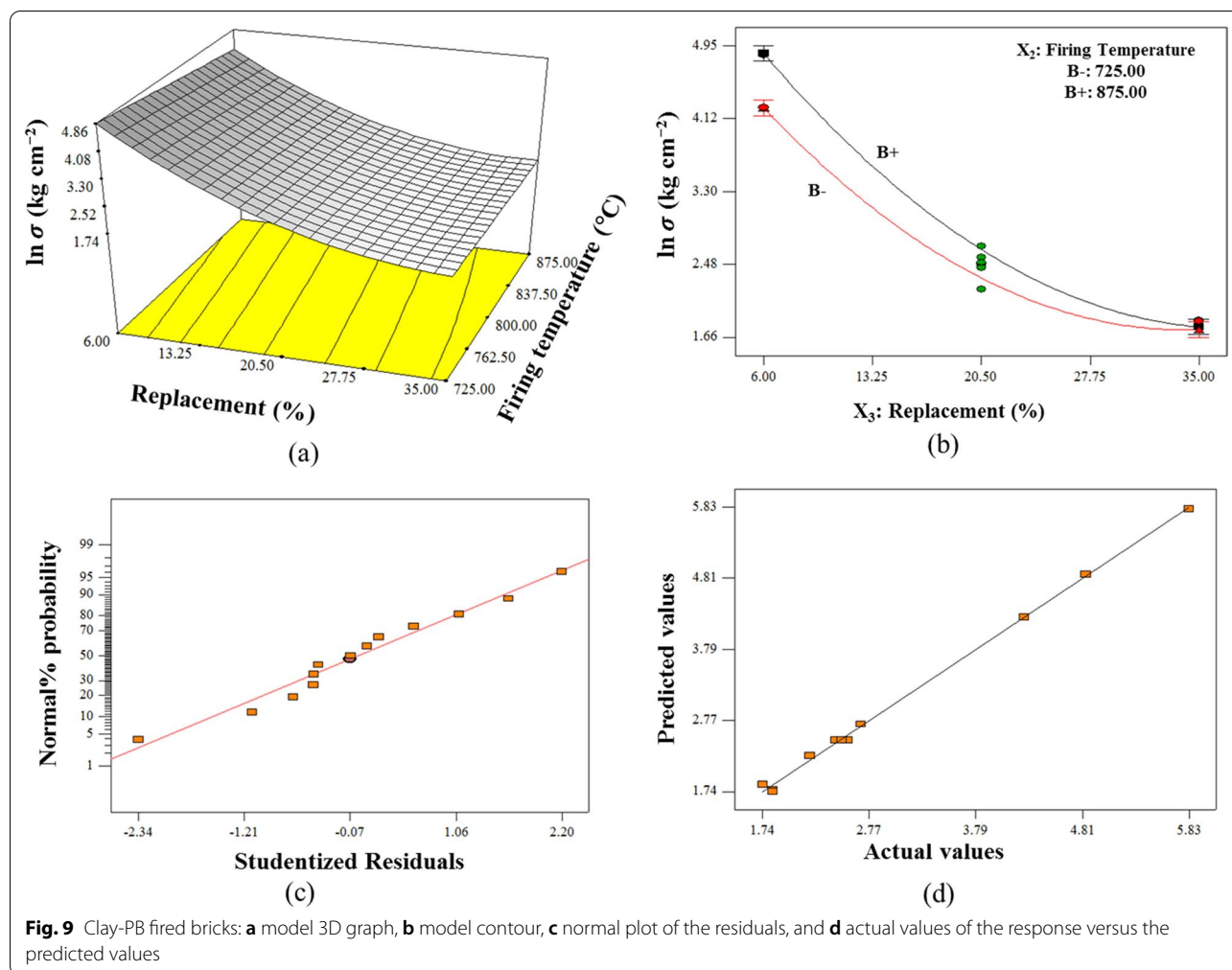
$$\text{TLS (\%)} = -4.72868 + 0.12579X_1 + 0.00690413X_2 \tag{11}$$

$$\text{CWA (\%)} = 13.87315 + 0.79680X_1 + 0.00759404X_1^2 \tag{12}$$

$$\text{BWA (\%)} = 15.66218 + 1.17997X_1 \tag{13}$$

$$\text{BD (g cm}^{-3}\text{)} = 1.77502 - 0.023388X_1 \tag{14}$$

As illustrated in Fig. 10, the model of LOI is a quadratic model that depends predominantly on the replacement percentage. The minimum LOI is achieved with a 6% clay replacement and 725 °C firing temperature. The TLS has a linear relationship between it and the investigated parameters. As can be seen in the regression equations, the clay replacement percentage has a dominant effect on TLS. The contour of TLS shows the minimum response is obtained for the minimum parameter values investigated.



**Table 11** Statistical analysis for the response

	Prediction	SE <sup>a</sup> mean	95% CI low	95% CI high	SE <sup>b</sup> Pred	95% PI low	95% PI high
ln(C. Strength)	4.859	0.054	4.74	4.98	0.088	4.66	5.06

<sup>a</sup> Standard error of the mean

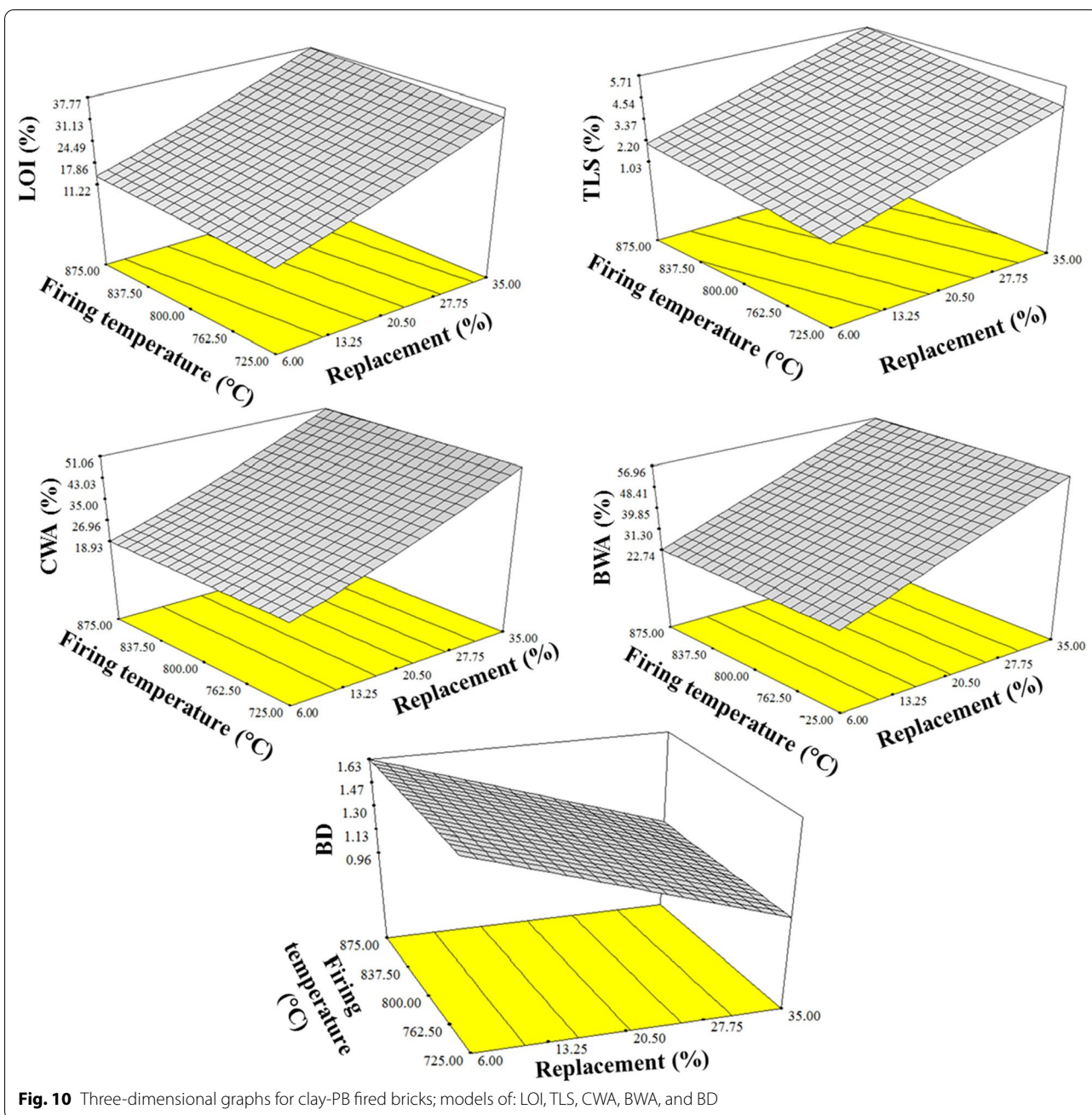
<sup>b</sup> Standard error of the prediction

The modeling of the remaining characteristics of clay-PB fired bricks shows considerable differences to clay-PA fired bricks. These variations are related to the material of the peanut shell. The modeling of the CWA is a quadratic model for only one parameter, and the firing temperature is not significant. The BWA has a linear relationship with the replacement percentage. The minimum values of both CWA and BWA were obtained with the minimum value of clay replacement. The BD also has a linear relationship with the clay replacement percentage, and the temperature is not a significant parameter. The lowest tested values of the parameters under investigation result

in a high BD and a sufficient SC of 0.89. Thus, the condition suggested by this investigation is that of a 6% clay replacement and a 725 °C firing temperature.

### Conclusions

Contaminated green waste represents a danger to human and animal health. It is the main cause of cancer and liver fibrosis, so the safe disposal of such waste is necessary unless propagation of the contamination of peanut crops can be prevented. This work showed that both the kernel and whole-grain peanut could be used as a clay replacement in the preparation of fired bricks to conserve



**Fig. 10** Three-dimensional graphs for clay-PB fired bricks; models of: LOI, TLS, CWA, BWA, and BD

natural resources and safely dispose of waste. The oil content of the peanut was found to decrease the required firing temperatures. Moreover, the fibrous material of the grain contributes to increasing the compressive strength of the green bricks. The RSM facilitates the optimization of the parameters affecting the quality of the product and creates models used to predict the characteristics of the product. Using a 6% clay replacement and a 725 °C firing temperature yields bricks of sufficient compressive strength (109.85 kg cm<sup>-2</sup> for clay bricks with PA and

126.33 kg cm<sup>-2</sup> for clay bricks with PB). This study paves the way for a more sustainable environment with zero-waste production.

**Abbreviations**

AFs: Aflatoxins; ANOVA: Analysis of Variance-Statistics; B1, B2, G1, G2: Mycotoxins types; BD: Bulk density; BWA: Boiling water absorption; CCD: Central Composite Design; CI: Confidence interval; CV: Coefficient of variance; CWA: Cold water absorption; DTA: Differential thermal analysis; EC: European Commission; FTIR: Fourier-transform infrared spectroscopy; IARC: International Agency for Research on Cancer; LOI: Loss on ignition; MPL: Maximum permissible

limit; P: Contaminated peanut; PA: Contaminated peanut kernel (without the outer shell); PB: Contaminated whole peanut grain (with the outer shell); RSM: Response surface method; TGA: Thermo-gravimetric analyses; TLS: Total linear shrinkage; SC: Saturation coefficient; XRD: X-ray diffraction; XRF: X-ray fluorescence.

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#### Author contributions

SAE-M performed the design of the experiments through response surface method, performed the computations, verified the analytical methods, and wrote the manuscript. ASS conceived of the presented idea and developed the theory, and performed the analysis of peanut feedstock. SKA conceived of the presented idea and developed the theory, performed the clay bricks experimental work, and wrote the manuscript. All authors discussed the results and contributed to the final manuscript. All authors read and approved the final manuscript.

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