



Bioconversion of Rice Straw as a Ruminant Feed Using Three Strains of White Rot Fungi

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Authors' contributions

This work was carried out in collaboration between both authors. Author AAW wrote the protocol, carried out the practical work, managed the analyses of the study, managed the literature and wrote the first manuscript while author JAA designed and supervised the study and also edited the first manuscript. Both authors read and approved the final manuscript.

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ABSTRACT

Aims: To determine nutritive value of rice straw samples biodegraded by different fungal strains.

Study Design: Completely Randomized Design.

Place and Duration of Study: Fungal inoculation of rice straw samples was carried out at the Federal Institute of Industrial Research Oshodi (FIIRO) Nigeria, for 21 days; proximate and fibre fractions analysis was done at the laboratory of the International Institute for Tropical Agriculture (IITA) Ibadan, Nigeria, while *in vitro* gas production was carried out at the laboratory of the Department of Animal Science, University of Benin, Nigeria.

Methodology: Samples of untreated rice straw (UTRS) were biodegraded for 21 days using three strains of edible mushroom *Pleurotus tuberregium* (PTTRS), *Pleurotus pulmonarius* (PPTRS) and *Pleurotus ostreatus* (POTRS). The substrates were analyzed for changes in the proximate composition and crude fibre fractions. *In vitro* Gas Production (IVGP) was used to predict the Metabolizable Energy (ME), Organic Matter Digestibility (OMD), Short Chain Fatty Acids (SCFA), methane production (CH₄), gas production at zero hour (a), total gas production (b), rate of fermentation (c) and incubation time (t).

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Results: Proximate composition showed crude protein (CP), ether extract (EE), ash and nitrogen free extracts (NFE) values to be 8.32, 15.32, 5.95 and 6.44%; 2.91, 3.93, 3.43 and 1.90%; 5.88, 12.38, 7.26 and 13.76% and 48.75, 43.65, 46.51 and 27.40% for UTRS, PTTRS, PPTRS and POTRS respectively. Crude fibre fractions for neutral detergent fibre (NDF), acid detergent fibre (ADF), acid detergent lignin (ADL), hemicellulose (HEM) and cellulose (CL) were 72.06, 77.43, 73.71 and 76.63%; 51.31, 6.14, 12.44 and 10.87%, 11.39, 1.43, 1.23 and 1.37%; 20.75, 71.29, 61.27 and 65.77% and 39.92, 4.71, 11.22 and 9.49% for UTRS, PTTRS, PPTRS and POTRS respectively. Estimates of ME, OMD, SCFA and CH₄ were 4.23, 7.29, 7.32 and 9.10 MJ/kg DM; 31.34, 52.94, 45.39 and 57.83%; 0.18, 0.56, 0.56 and 0.78 $\mu\text{mol}/200 \text{ mg DM}$ and 13.50, 9.33, 10.33 and 10.67 ml for UTRS, PTTRS, PPTRS and POTRS respectively. IVGP characteristics for a, a+b, b, c and t were 1.25, 2.38, 1.13 and 1.57; 9.25, 23.88, 24.13 and 24.07; 8.00, 21.50, 23.00 and 22.50; 0.11, 0.41, 0.35 and 0.35 and 21.00, 8.00, 9.00 and 9.00 hours for UTRS, PTTRS, PPTRS and POTRS respectively

Conclusion: Based on the result, *Pleurotus tuberregium* degraded rice straw (PTTRS) was adjudged superior to the others due to the higher crude protein, ether extract, potentially degradable fractions, high gas production per unit time and lower methane production.

Keywords: *In vitro*; nutrients; rice straw; degraded; mushroom.

1. INTRODUCTION

Huge amounts of agricultural waste are produced from crop farms all over the world annually [1], one of which is rice straw. Rice straw, a residue arising from rice farming, is produced in copious quantities worldwide annually. It is a residue of the rice plant left after the grains have been harvested [2]. It is reported to be low in nitrogen, rich in polysaccharides and has a high lignin and silica content, limiting voluntary intake and reducing degradability by ruminal micro organisms all of which serve as a hindrance to its effective utility as a livestock feed [3,2].

Biodegradation of crop residues using edible mushroom has been reported to give rise to products with enhanced nutritive value that promote farm animal performance [4]. Thus rice straw could also be converted to a useful animal feed resource through biological treatment. This is particularly useful to ensure provision of animal protein sources for the teeming world population. Animal nutritionists are continuously evaluating forages and other feed resources so as to provide solutions to problems arising from practical feeding of livestock in specific situations and on a year round basis. The *in vitro* gas production technique of assessing nutritive value of feeds and feedstuffs [5] is particularly convenient due to less requirement of time, less cost and ability to analyze many samples almost simultaneously. The use of chemical composition in combination with *in vitro* digestibility and degradability can be a useful tool for evaluation of nutritive value of feed resources [6]. It was

suggested [7] that the *in vitro* gas technique is precise in measuring both rate and extent of fermentation in addition to the residue of fermentation at various times of incubation. The method had been used to assess the replacement value of forages [8], analyze forages [6] and to determine the nutritive value of differently processed feed inputs [9]. The aim of this study was to evaluate rice straw samples ensuing from biodegradation by three edible mushrooms *Pleurotus tuberregium*, *Pleurotus pulmonarius* and *Pleurotus ostreatus* by assessing the proximate composition, crude fibre fractions, *in vitro* gas production and nutrient estimations.

2. MATERIALS AND METHODS

2.1 Sample Collection

Samples of dried rice straw were collected from crop farms around the environs of the University of Agriculture, Makurdi, Benue State. The samples were milled and oven dried at 65°C until a constant weight was obtained for any dry matter determination.

2.2 The Fungi

The sporophores of *Pleurotus tuber-regium*, *Pleurotus pulmonarius* and *Pleurotus ostreatus* growing in the wild were collected from University of Ibadan Botanical garden. These were tissue cultured to obtain fungal mycelia [10]. The pure culture obtained was maintained on plate of potato dextrose agar (PDA).

2.3 Degradation of Rice Straw by Different Fungi

2.3.1 Preparation of substrates

The jam bottles used for this study were thoroughly washed, dried for 10 min. at 100°C. 25.00 g of the dried milled substrate was weighed individually into each jam bottle and 70 ml distilled water was added. The bottle was immediately covered with tin foil and sterilized at 121°C for 15 min. Each treatment was done in triplicate.

2.3.2 Inoculation

Each bottle was inoculated at the centre of the substrate with two, 10.00 mm mycelia disc and covered immediately. They were kept in the dark cupboard in the laboratory at 30°C and 100% relative humidity (RH). At day 21 of inoculation, the experimental bottles were autoclaved to terminate the mycelia growth. Samples of biodegradations were oven dried to constant weight for chemical analysis and *in vitro* digestibility.

2.3.3 *In vitro* gas production

This was carried out at the Animal Science laboratory of University of Benin. Rumen fluid was obtained from three West African Dwarf male goats through suction tube via the oesophagus before morning feed. The animals were fed with 40% concentrate (40% corn, 10% wheat offal, 10% palm kernel cake, 20% groundnut cake, 5% soybean meal, 10% brewers grain, 1% common salt, 3.75% oyster shell and 0.25% fishmeal) and 60% Guinea grass. Incubation was carried out [5] in 120 ml calibrated syringes in three batches at 39°C. To 200 mg sample in the syringe was added 30 ml inoculum containing cheese cloth strained rumen liquor and buffer (9.8 g NaHCO₃ + 2.77 g Na₂HPO₄ + 0.57 g KCL + 0.47 g NaCl + 0.12 g MgSO₄.7H₂O + 0.16 g CaCl₂. 2H₂O in a ratio (1:4 v/v) under continuous flushing with CO₂. The gas production was measured at 3, 6, 9, 12, 15, 18, 21, and 24 hrs. After 24 hr of incubation, 4 ml of NaOH (10M) was introduced to estimate the amount of methane produced [11]. The average volume of gas produced from the blanks was deducted from the total volume of gas produced. Fermentation characteristics were estimated using the equation $Y = a + b(1 - e^{-ct})$ [12], where Y = volume of gas produced at time 't', a = intercept (gas produced from the soluble

fraction), b = gas production rate constant for the insoluble fraction, (a + b) = final gas produced, C = gas production rate constant for the insoluble fraction (b), t = incubation time. The average volume of gas produced from the blanks was deducted from the total volume of gas produced. Gas production was measured using the indices a= gas production at zero hour, b= total gas production and t= incubation time. Metabolizable energy (ME, MJ/Kg DM) and organic matter digestibility (OMD %) were estimated [5] and short chain fatty acids (SCFA) was calculated [13]. $ME \text{ MJ/kg DM} = 2.20 + 0.136 * Gv + 0.057 * 0.0029 CF$; $OMD = 14.88 + 0.88Gv + 0.45CP + 0.651XA$; $SCFA = 0.0239 * Gv - 0.0601$; Where Gv, CP, CF and XA are net gas production (ml /200 mg DM), crude protein, crude fibre and ash of the incubated sample respectively.

2.4 Chemical Composition

Dry matter was obtained by oven drying the milled samples to a constant weight at 105°C for eight hours. Crude protein was determined as Kjeldhal nitrogen x 6.25 ether extracts, crude fibre and ash were determined according to [14] method. Neutral detergent (NDF), Acid detergent fibre (ADF) and Acid Detergent Lignin (ADL) were determined using the method described by [15].

Hemicellulose was calculated as the difference between NDF and ADF while cellulose was calculated as the difference between ADF and ADL.

2.5 Statistical Analysis

Data obtained were subjected to Analysis of Variance (ANOVA) using the Minitab Statistical software which also separated the means where significant differences occurred (at 95% probability).

3. RESULTS

3.1 Proximate Composition and Crude Fibre Fractions

The proximate composition and crude fibre fractions of rice straw treated with different fungal species is shown in Table 1. There was a significant (P = 0.01) increase in the crude protein (CP) of the fungal treated RS compared to the untreated RS. The highest CP was by the PTTTS, followed by POTRS while CP of PPTRS

was lowest. Ether extract was also significantly ($P = 0.01$) affected, being highest in PTTRS, followed by PPTRS and least in POTRS. Ash percent was significantly ($P = 0.00$) highest in POTRS, followed by PTTRS and lowest in PPTRS, while the nitrogen free extracts fraction was significantly ($P = 0.03$) lowest in POTRS, followed by PTTRS and then PPTRS.

Neutral detergent fibre (NDF) was significantly ($P = 0.03$) highest in PTTRS, followed by POTRS and then PPTRS. Acid detergent fibre (ADF) values were significantly ($P = 0.00$) reduced in the order PPTRS, POTRS and then, PTTRS. Acid detergent lignin (ADL) percent was significantly ($P = 0.00$) least in PPTRS, followed by POTRS and lastly PTTRS. Hemicellulose percent was significantly ($P = 0.00$) highest in PTTRS, followed by POTRS and PPTRS, while cellulose percent was significantly ($P = 0.02$) least in PTTRS, followed by POTRS and PPTRS.

3.2 Nutrient Estimations

The Metabolizable energy (ME), organic matter digestibility (OMD), short chain fatty acid (SCFA) and Methane production are shown in Table 3.

The ME was significantly ($P = 0.00$) improved, being highest in POTRS while values of PTTRS and PPTRS were similar to each other. The OMD was significantly ($P = 0.04$) increased, being highest in POTRS, followed by PTTRS and then PPTRS. The SCFA values were also significantly ($P = 0.04$) improved, following pattern exhibited by ME. Methane production was significantly ($P = 0.01$) reduced due to fungal treatment, being lowest in PTTRS, followed by PPTRS and then POTRS.

3.3 In vitro Gas Production

The *in vitro* gas production characteristics are shown in Table 3. Gas production at zero hour (a) was not significantly ($P = 0.02$) affected by fungal treatment. Total gas production (b) was significantly ($P = 0.02$) affected by fungal treatment, being highest in PPTRS and POTRS, which were similar to each other, but higher than PTTRS which was also significantly higher than the UTRS. Gas production rate (c) was significantly ($P = 0.01$) highest in PTTRS, followed by PPTRS and POTRS which were similar to each other, but significantly different from UTRS. Incubation time (t) was significantly

Table 1. Proximate composition and crude fibre fractions of rice straw degraded by different fungal species

Parameter	UTRS	PTTRS	PPTRS	POTRS	SEM
Crude protein (%)	8.32 ^b	15.32 ^a	5.95 ^d	6.44 ^c	0.11
Ether extract (%)	2.91 ^c	3.93 ^a	3.43 ^b	1.90 ^d	0.09
Ash (%)	5.88 ^d	12.38 ^b	7.26 ^c	13.76 ^a	0.24
Nitrogen free extracts (%)	48.75 ^a	43.65 ^c	46.51 ^b	27.40 ^d	0.63
NDF	72.06 ^d	77.43 ^a	73.71 ^c	76.63 ^b	0.02
ADF	51.31 ^a	6.14 ^d	12.44 ^b	10.87 ^c	1.20
ADL	11.39 ^a	1.43 ^b	1.23 ^d	1.37 ^c	1.35
HEM	20.75 ^d	71.29 ^a	61.27 ^c	65.77 ^b	0.03
CL	39.92 ^a	4.71 ^d	11.22 ^b	9.49 ^c	2.31

^{a,b,c,d} = Means in same row with different superscripts vary significantly ($P=0.05$)

UTRS = Untreated rice straw; PTTRS = *Pleurotus tuberregium* treated rice straw; PPTRS = *Pleurotus pulmonarius* treated rice straw; POTRS = *Pleurotus ostreatus* treated rice straw; NDF = Neutral detergent fiber; ADF = Acid detergent fiber; ADL = Acid detergent lignin; HEM = Hemicellulose; CL = Cellulose

Table 2. Metabolizable energy, organic matter digestibility, short chain fatty acids and methane production of rice straw degraded by different fungal species

Treatment	ME (MJ/kg DM)	OMD (%)	SCFA (umol/200 mg DM)	Methane (g/100 ml)
UTRS	4.23 ^c	31.34 ^d	0.18 ^c	13.50 ^a
PTTRS	7.29 ^b	52.94 ^b	0.56 ^b	9.33 ^d
PPTRS	7.32 ^b	45.39 ^c	0.56 ^b	10.33 ^c
POTRS	9.10 ^a	57.83 ^a	0.78 ^a	10.67 ^b
SEM	0.13	0.51	0.05	0.20

^{a,b,c,d} = Means in same column with different superscripts vary significantly ($P<0.05$)

UTRS = Untreated rice straw; PTTRS = *Pleurotus tuberregium* treated rice straw; PPTRS = *Pleurotus pulmonarius* treated rice straw; POTRS = *Pleurotus ostreatus* treated rice straw; ME = Metabolizable energy; OMD = Organic matter digestibility; SCFA = Short chain fatty acids

Table 3. *In vitro* gas production characteristics of rice straw degraded by different fungal species (ml/200 mgDM)

Fermentation characteristics	UTRS	PTTRS	PPTRS	POTRS	SEM
A	1.25	2.38	1.13	1.57	0.44
B	9.25 ^b	23.88 ^a	24.13 ^a	24.07 ^a	0.78
C	0.11 ^c	0.41 ^a	0.35 ^b	0.35 ^b	0.78
T	21.00 ^a	8.00 ^c	9.00 ^b	9.00 ^b	0.50

^{a,b,c,d} = Means in same row with different superscripts vary significantly ($P < 0.05$)

UTRS = Untreated rice straw; PTTRS = *Pleurotus tuberregium* treated rice straw;

PPTRS = *Pleurotus pulmonarius* treated rice straw; POTRS = *Pleurotus ostreatus* treated rice straw;

a = gas production at zero hour; b = total gas production; c = Rate of fermentation;

t = incubation time

($P = 0.00$) depressed by fungal treatment, being lowest in PTTRS, followed by PPTRS and POTRS which were similar to each other, but significantly lower than UTRS.

4. DISCUSSION

4.1 Proximate Composition and Fibre Fractions

The highest CP level of PTTRS implies that the fungus effectively degraded the rice straw and colonized it causing the CP level to rise due to CP of the fungal biomass and enzymes. According to [16], increase in CP levels of fungal treated materials is caused by secretion of certain extra cellular enzymes which are proteinous in nature during the materials breakdown and its subsequent metabolism. That the CP of PPTRS and POTRS was lower than that of UTRS implies that the fungi were less effective in degrading rice straw. The highest EE percent of PTTRS and PPTRS means the fungi contributed more EE while in POTRS; it used EE as a nutrient in its development. According [17], some micro organisms consume some fatty acids as suitable energy for growth. The general increment in ash values of the treated straw means more minerals and vitamins were available after the degradation. This observation is consistent with the report of [10] who reported improved values for ash when maize husk was degraded using different species of white rot fungi. The Lower nitrogen free extracts fraction in all the treated rice straw means that the fungi used carbohydrate as source of energy.

NDF and hemicellulose fractions generally increased due to fungal treatment means the fungi secreted enzymes less effective in breaking down the components. It was reported [18] that hemicellulose percent was increased when rice

straw was treated with *Trichoderma reesi*. The generally reduced levels of ADF, ADL and cellulose demonstrate the fact that the fungi were more effective in degrading those fractions in the straw.

4.2 Nutrient Estimation

The estimated Metabolizable energy increased signifying that fungal treatment produced more nutritive products. This result is not unexpected because ME estimation makes use of gas volume, protein, fat and crude fibre fraction of the feeds.

Organic matter digestibility also improved due to fungal treatment. This is reasoned to be due to reduction in lignin which inhibits digestibility and the high ash value and gas volume used in estimating the organic matter digestibility.

Short chain fatty acid production increase in the treated materials is reasoned to be due to higher absolute gas production, which was most evident in the 24 hour of incubation. Close association between SCFA and gas production *in vitro* was reported [14]. During incubation of feedstuff with buffered rumen fluid *in vitro*, carbohydrates are fermented to SCFA and gases, mainly carbon dioxide and methane and microbial cells. This report implies the release of more energy courtesy of fungal treatment.

Methane production decreased due to fungal treatment. Generally, methane represents source of energy waste or loss, as the animals have to eructate or belch it to prevent bloat [19]. This reduction in methane production makes the treated materials more environmentally friendly, as they would contribute less to global warming capable of frustrating food production in agricultural areas due to drought or flood [20].

4.3 *In vitro* Gas Production

Gas produced at zero hour (a) did not significantly differ, but was higher in PTTRS showing its tendency to release more energy from the soluble fractions than the other materials. The total gas produced (b) increased in the treated materials. This result is consistent with those of other workers [10,21]. The improved rate of gas production (c) in fungal treated materials signified more carbohydrate availability per unit time and also how fast the treated materials would release nutrients for animal use. It had been reported that rate of gas production determines digestion time and consequently how long a potentially digestible material would occupy space [22]. Thus, the higher gas production rate of the treated materials would make them to be easily digested and other materials fed consequently leading to better performance by animals fed the materials. In general, gas production can be regarded as an indicator of carbohydrate degradation and the low gas production in the UTRS could be explained by lignin binding to the carbohydrate [23]. On the other hand higher gas production of the treated straw is further explained by higher CP (especially PTTRS) and the generally reduced lignin. According to [24], lignin protects carbohydrates from attack by rumen microbes and negatively influences *in vitro* gas production. Gas production on incubation of feeds in buffered rumen fluid is associated with feed fermentation and carbohydrate because gas production from protein fermentation is relatively small as compared to carbohydrate fermentation while contribution of fat to gas production is negligible [20].

The reduced incubation time (t) of the treated materials implies that it takes less time for nutrients to be released. This would be beneficial as the animals will be able to tap nutrients in the materials faster, not having to wait for long before benefiting from feed intake.

5. CONCLUSION

It was concluded that treatment of rice straw with *Pleurotus tuberregium* significantly improved its nutritive value as there was increase in crude protein and gross energy, while crude fibre and fibre fractions decreased. Also, the fungal treatment of rice straw significantly improved *in vitro* gas production and Metabolizable energy, organic matter digestibility and short chain fatty acids. Additionally, methane production was significantly decreased by fungal treatment of the

rice straw, thereby making the resultant product more environmentally friendly and of less potential in contributing to global warming.

6. RECOMMENDATION

It was recommended that fungal treatment of rice straw before feeding to animals should be encouraged due to the additional benefit of reduced methane production.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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