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Biomass Residues as Power Station Fuels

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Introduction



Feedstock for co-firing in the UK by type, quantity and source

	Feedstock	Quantity burned (tonnes) In 2005	% quantity burned (tonnes) In 2005	Likely country of origin	Mode of transport	Total transport- related emissions (kg CO ₂ /tonne biomass)
	Energy crops (SRC,granulated willow, miscanthus)	4,306	0.3	UK	Road	1.7
+	Shea residues (meal and pellets)	5,420	0.4	Africa	Ship	55.4
	Sunflower pellets	20,331	1.4	Romania	Road & ship	47.1
	Sewage sludge and waste derived fuels	49,155	3.5	υκ	Road	3.4
	Cereal co products and pellets	102,246	7.2	UK	Road	1.7
	Tallow	119,828	85	UK	Road	1.7
+	 Olive waste (residue and expeller) 	283,222	20.1	Greece, Italy Spain	Road & ship	21.2
	Wood (sawdust, chips, pellets, tall oil)	377,956	26.8	UK, Canada, Latvia, Scandinavia	Road & ship	1.7 (UK) to 42.9
	 Palm residues (palm kernel expeller, shell, pellets, oil) 	449,657	31.8	Indonesia, Malaysia	Road & ship	106.5 (Indonesia) to 107.4 (Malaysia)
	Total mass	1,412,121				
	Total energy (PJ)	14.1				

Sources: UK Biomass Strategy, DEFRA, May 2007 & Evaluating the Sustainability of Co-firing in the UK, report to DTI from Themba Technology Ltd, September 2006

Introduction: Imported biomass



Palm kernel expeller Shea residue Olive residue



- Oil extracted from both palm fruit (flesh) and kernel (nut)
- PKE: fibrous remains from the kernel oil extraction process.

- Shea butter extracted from kernel of shea fruit
- Residue: fleshy

mesocarp, shell and husk left after removal of butter

• Olive residues: crushed olive kernel, shell, pulp, skin

• Imported as cake, expeller, or pellets

Fuel samples provided by RWE nPower

Aims



To examine the combustion properties of PKE, shea residue, and three different olive residues

To examine the nitrogen partitioning and char-N chemistry during combustion

Methodology



- Fuel characterisation: ultimate, proximate, and metal analyses, ash behaviour (fouling and slagging indices)
- Nitrogen partitioning
- Studies of fuel chars:
 - 1. Characterisation: ultimate and proximate analyses
 - 2. Combustion by TGA-MS: char-N conversion to Ncontaining species

Results: Fuel characterisation



Parameter	PKE	Shea residue	Olive residue A	Olive residue B	Olive residue C
C (% daf)	51.12	54.24	54.42	54.33	51.38
H (% daf)	7.37	6.58	6.82	7.20	6.32
N (% daf)	2.80	3.48	1.40	1.39	1.45
O ($\%$ daf) ^a	38.71	35.70	37.36	37.08	40.85
C/N	21.32	18.21	45.41	45.59	41.33
Moisture (% ar)	7.60	8.42	6.40	4.61	5.19
Volatiles (% ar)	72.12	57.06	65.13	70.68	55.51
Fixed carbon $(\% \text{ ar})^a$	16.18	27.62	19.27	17.17	17.31
Ash (% ar)	4.10	6.90	9.20	7.54	21.99
HHV (MJ/kg) dry basis ^b	20.00	20.37	19.67	20.25	16.10
Ash composition (% dry ba	asis)				
Al_2O_3	0.87	1.29	1.94	0.85	2.74
CaO	11.90	5.51	15.44	9.40	19.49
Fe ₂ O ₃	5.70	2.37	2.14	0.75	5.29
K ₂ O	21.43	42.57	31.04	32.08	4.41
MgO	11.51	6.83	5.78	2.87	5.25
Mn ₃ O ₄	1.03	0.05	0.05	0.02	0.33
Na ₂ O	0.41	0.95	0.47	0.33	0.35
SiO ₂	16.51	14.40	21.10	10.88	67.40
Total ash components	69.35	73.97	77.96	57.18	105.25

^a calculated by difference, ^b calculated by method in Friedl et al. 2005

Results: Slagging and fouling indices



Fuel	Alkali index	Base to	Base	
	(kg alkali/GJ)	acid ratio*	percentage	
PKE	0.48	2.93	50.94	
Shea residue	1.61	3.71	58.23	
Olive residue A	1.57	2.38	54.87	
Olive residue B	1.27	3.88	45.44	
Olive residue C	0.69	0.50	34.79	

*TiO₂ not included

(Jenkins et al. 1998)

$$R_{b/a} = \% (\underline{Fe_2O_3 + CaO + MgO + K_2O + Na_2O}) \\ \% (SiO_2 + TiO_2 + Al_2O_3)$$

Al>0.34 kg alkali/GJ \longrightarrow fouling virtually certain! (Miles et al. 1996)

Biomass ash softening temperatures





Results: Fuel chars and nitrogen partitioning



Parameters	PKE	Shea residue	Olive residue A	Olive residue B	Olive residue C
C (% daf)	91.46	89.34	84.30	85.78	86.23
H (% daf)	2.74	3.14	2.50	2.64	3.48
N (% daf)	4.37	2.49	1.10	1.40	1.18
O $(\% \text{ daf})^a$	1.43	5.04	12.10	10.17	9.12
C/N	24.41	41.92	89.48	71.49	85.57
Moisture (%) ^b	0.26	0.98	0.00	0.81	0.36
Ash (% dry basis) ^b	62.44	32.20	36.78	40.90	73.65
Char yield (% dry basis) ^c	14.76	39.59	26.95	33.06	44.38
Volatile yield (% dry basis) ^c	85.24	60.41	73.05	66.94	55.62
N partitioning					
N (%) in char	9.03	20.67	18.22	17.32	12.22
N (%) in volatiles	90.97	79.33	81.78	82.68	87.78

^a calculated by difference

^b from combustion in STA-MS (hr 10 ℃ min⁻¹ to 600 ℃)

° from char preparation (hr 10 ℃ ms⁻¹ to 1000 ℃)

Results: Char-N conversions



Masses monitored: $m/z \ 14: N_2^{2+} and CO^{2+}$ $m/z \ 27: HCN + tail end of$ $m/z \ 28 signal$ $m/z \ 28:^{12}C^{16}O$ $m/z \ 30: NO + {}^{12}C^{18}O$ $m/z \ 43: HCNO$ $m/z \ 44: {}^{12}C^{16}O_2 + N_2O$ $m/z \ 46: NO_2 + {}^{12}C^{18}O^{16}O$ $m/z \ 52: C_2N_2$



PKE char

Shea residue char



Results: Char-N conversions



Olive residue A char

Olive residue B char

Olive residue C char









Results: Char-N conversions



 * from $N_2{}^{2+}$ signal and m/z 14:m/z 28 ratio=0.154

Conclusions



- All fuels investigated show a high tendency to fouling and slagging, with ash softening temperatures <1100 ℃
- Most of the N is lost in the volatiles (~80-90%)
- Most of the char-N is released as N_2 and NO_x in combustion
- Other species detected: HCN and C₂N₂ in small quantities (from PKE and olive residue B chars)
- Similarly to coal, N is retained until near the end of char combustion stage, with NO as a primary product

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