



AN OVERVIEW OF THE CHEMICAL COMPOSITION OF BIOMASS*

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Introduction and Scope



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JRC-IE is conducting a detailed review of the biomass literature (>370 references) plus some key investigations related to biomass use in bioenergy with two fundamental objectives:

(1) to extend and improve the basic knowledge on composition and properties
(2) to apply this knowledge for the most advanced and environmentally safe utilization

Increased use of biomass for bioenergy will environmental impact, but what might be that impact?

Question: Can methodology for characterisation of coal be applied to biomass?

Question: What is an appropriate methodology for full characterisation of biomass for bioenergy, particularly co-firing applications?



Basic Methodology



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Initial Step:

establish the bulk chemical composition of biomass:

traditionally this should include -

proximate analysis (FC, VM, A, M),

ultimate analysis (C, O, H, S, N),

ash analysis (Si, Al, Fe, Ca, S, Mg, K, Ti, Na and P oxides)

Unfortunately, complete datasets for many varieties of biomass are quite limited (only for 86 varieties of biomass could be used for the present study after consideration of 280 references)



Aims of the Work



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This includes:

- evaluating the chemical composition of 86 varieties of biomass and their ashes based on the 19 parameters, plus additional data for Cl and Mn
- conducting some chemical comparisons between biomass and 38 Solid Fossil Fuels (SFF)
- establishing some basic trends/relationships based on biomass chemical analysis data
- proposing possible initial classifications and potential applications of the results



General classification of biomass



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According to origin

Biomass group	Biomass sub-groups, varieties and species		
1. Wood and woody biomass	Coniferous or deciduous; soft or hard; stems, branches, foliage, bark, chips, pellets, briquettes, sawdust, sawmill and others from various wood species		
2. Herbaceous and agricultural biomass	Annual or perennial and field-based or processed-based such as: 2.1. Grasses and flowers (alfalfa, arundo, bamboo, bana, brassica, cane, miscanthus, switchgrass, timothy) 2.2. Straws (barley, flax, oat, rape, rice, rye, wheat) 2.3. Other residues (fruits, shells, husks, pits, seeds, coir, stalks, cobs, bagasse, food, fodder, pulps)		
3. Aquatic biomass	Marine or freshwater algae; macroalgae or microalgae; blue, green, blue- green, brown, red, seaweed, kelp, lake weed, water hyacinth		
4. Animal and human biomass wastes	Bones, meat-bone meal, chicken litter, various manures		
5. Contaminated biomass and industrial biomass wastes (semi-biomass)	Municipal solid waste, demolition wood, refuse-derived fuel, sewage sludge, hospital waste, paper-pulp sludge, waste papers, paperboard waste		
6. Biomass mixtures	Blends from the above varieties		



Phase composition of biomass



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Matter	State and type	Phases and components
Organic matter	Solid, non-crystalline	Structural ingredients, mainly cellulose, hemicellulose, lignin
	Solid, crystalline	Organic minerals such as Ca-Mg-K-Na oxalates, others
Inorganic matter	Solid, crystalline	Mineral species from phosphates, carbonates, silicates, chlorides, sulphates, oxyhydroxides, nitrates, and other mineral classes
	Solid, semi-crystalline	Poorly crystallized forms of some silicates, phosphates, hydroxides, others
	Solid, amorphous	Amorphous phases such as various glasses, silicates, others
Fluid matter	Fluid, liquid, gas	Moisture, gas and gas-liquid inclusions associated with both organic and inorganic matter



Origin of Phases in Biomass



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Process	Place	Time	Formation mechanism
Natural	Authigenic (formed in	Syngenetic (during plant growing)	Generated by biogenic processes
	biomass)	Epigenetic (after plant died)	Originated by natural processes
	Detrital (formed outside biomass, but fixed in/on biomass)	Pre-syngenetic (before plant growing)	Pre-existing and finely dispersed mineral grains (commonly <1 μ m) introduced into the plant by water suspensions during syngenesis
		Pre-syngenetic, syngenetic or epigenetic	Pre-existing and fine-grained particles (normally <10-100 μm) introduced by water and wind on plant surfaces
Anthropogenic	Technogenic (formed outside or inside biomass and fixed in/on biomass)	Post-epigenetic (during and after plant collecting)	Natural and/or industrial components introduced in biomass during collecting, handling, transport and subsequent processing



Recalculating Chemical Composition



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Proximate and ultimate analyses data recalculated on <u>dry and</u> <u>daf basis</u>, respectively, results in different enrichment - depletion trends:

In decreasing order of abundance, the elements in biomass are normally:

- (1) major (>1.0%): C, O, H, N, Ca, K
- (2) minor (0.1-1.0%): Si, Mg, Al, S, Fe, P, Cl, Na
- (3) trace elements (<0.1%): Mn, Ti and others

Biomass group and sub-group	Enriched in	Depleted in
1. Wood and woody biomass (WWB)	Ca, Mg, Mn, VM	A, Cl, N, P, S, Si
2. Herbaceous and agricultural biomass (HAB)	FC, K, O, VM	C, H, Ca
2.1. Grasses (HAG)	K, O, Si, VM	Al, C, Ca, H, Na
2.2. Straws (HAS)	Cl, K, O, Si	C, H, Na
2.3. Other residues (HAR)	FC, K, Mg, P	Cl
3. Animal biomass (AB)	A, C, Ca, Cl, H, N, Na, P, S	Al, Fe, M, Mg, Mn, O, Si, Ti, VM
4. Contaminated biomass (CB)	A, Al, C, Cl, Fe, H, N, S, Ti	FC, K, P

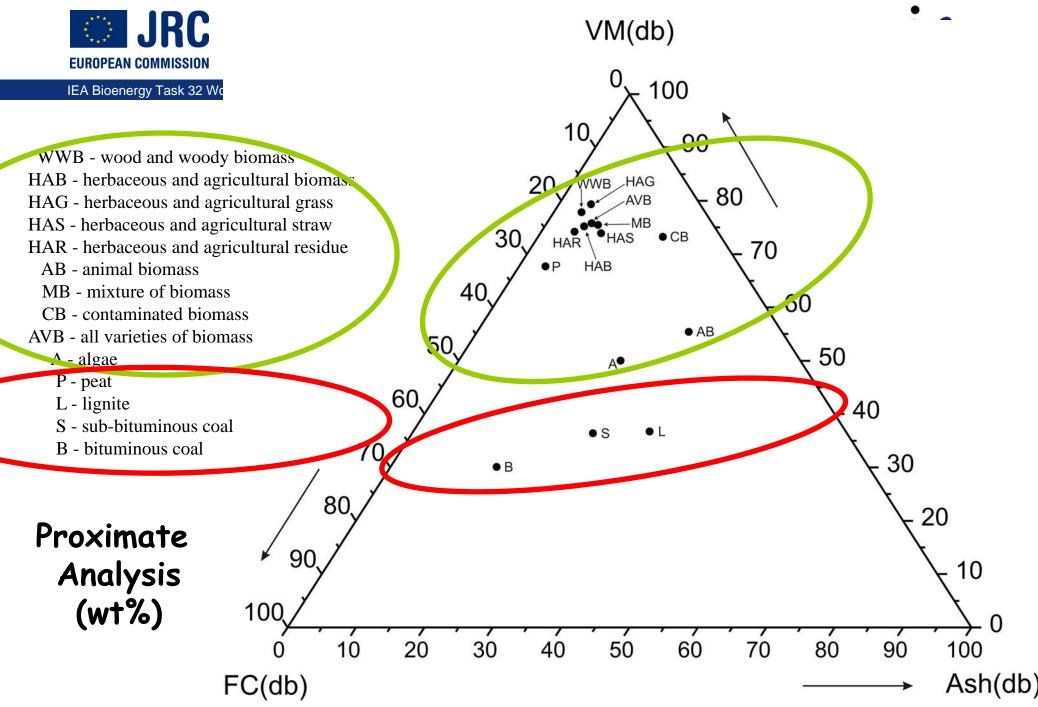


Intermediate Conclusions



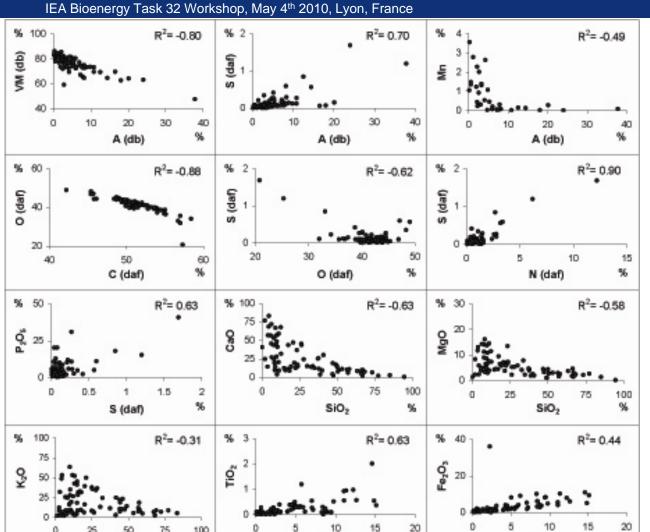
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- Chemical composition of natural biomass is simpler than that of Solid Fossil Fuels
- However, processing results in incorporation of numerous non-biomass materials
- Biomass composition is significantly different from that of SFFs
- The variations among biomass composition are greater than for coal. In comparison with coal, natural/virgin biomass (78 varieties) commonly shows:
 - (1) enrichment in Mn > K > P > Cl > Ca > (Mg, Na) > O
 - > VM > H
 - (2) depletion in S > (A, Al, Ti) > FC > (Fe, Si) > C > N





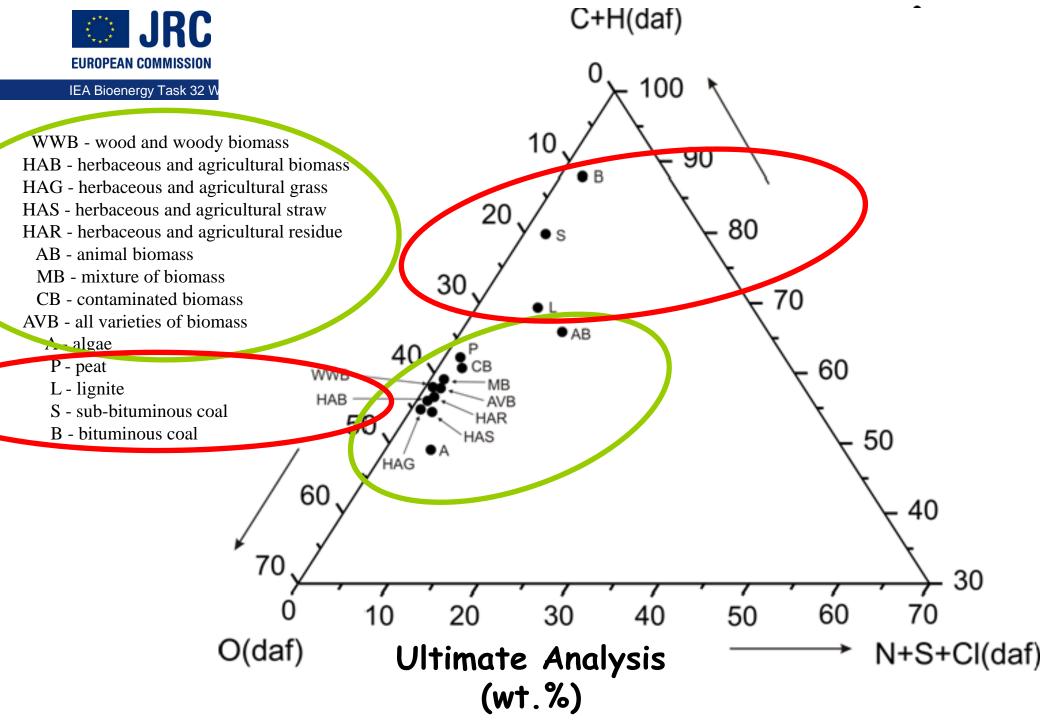




Five strong correlations for natural/biomass system (78 varieties):

- (1) C H
- (2) N S CI
- (3) Si Al Fe Na Ti
- (4) Ca Mg Mn
- (5) K P S Cl

Fig. 2. Some significant correlations among the chemical composition of biomass (wt.%)



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SiO₂+Al₂O₃+Fe₂O₃+Na₂O+TiO₂

biomass ash:

(1) $SiO_2 + Al_2O_3 + Fe_2O_3 +$ $Na_2O + TiO_2$

$$(2)$$
 CaO + MgO + MnO

(3) $K_2O + P_2O_5 + SO_3 +$

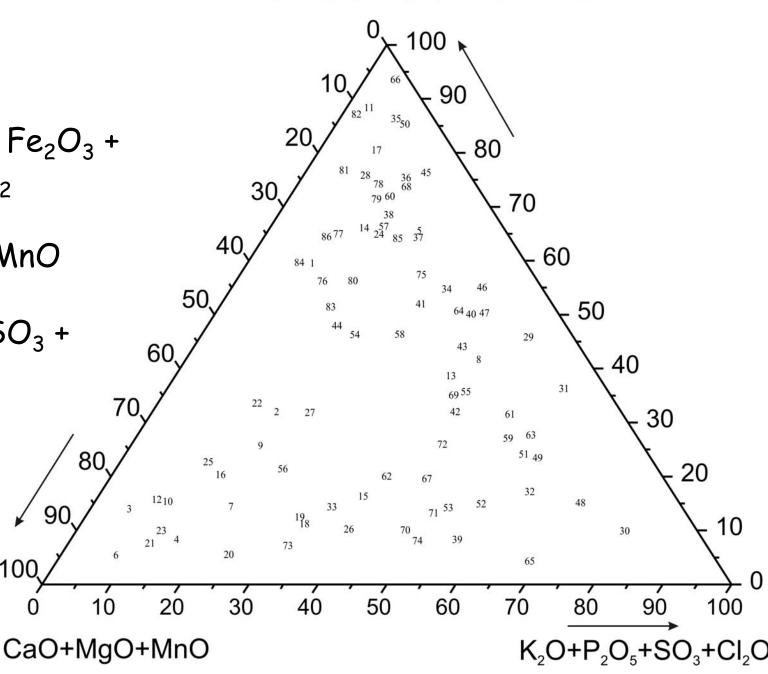
70,

80,

10

90

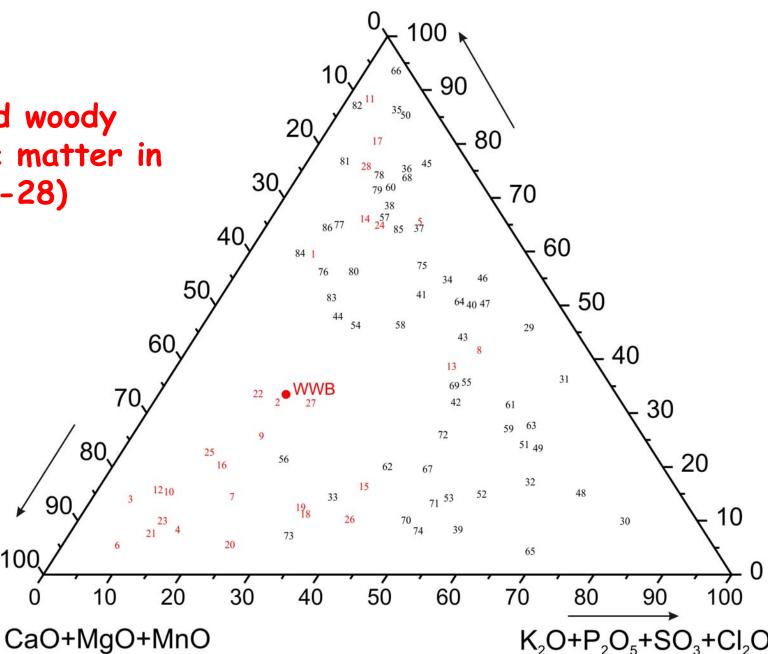
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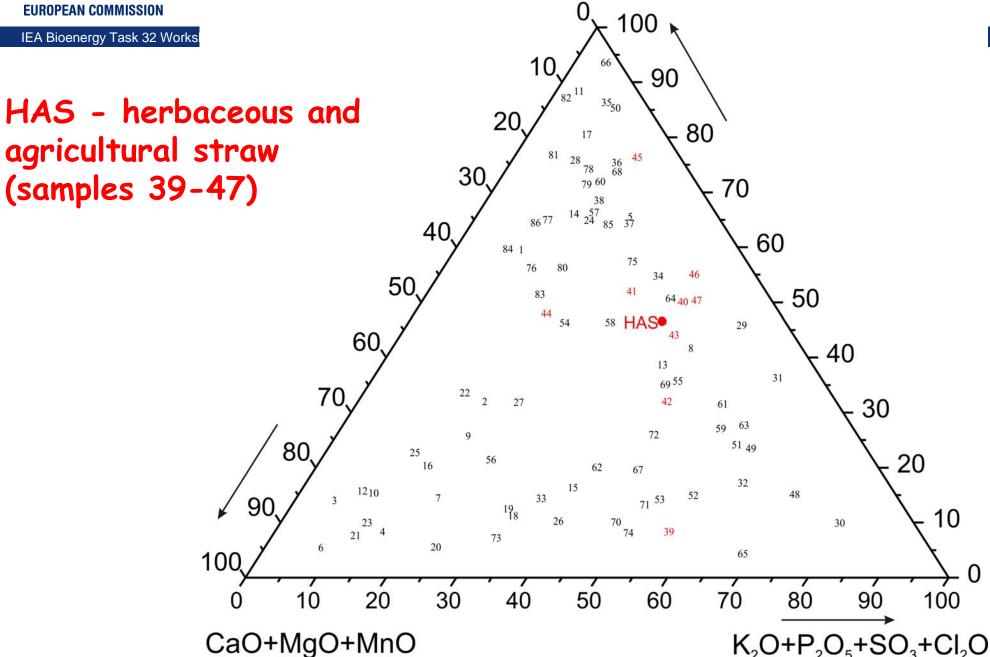


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WWB - wood and woody biomass inorganic matter in ashes (samples 1-28)

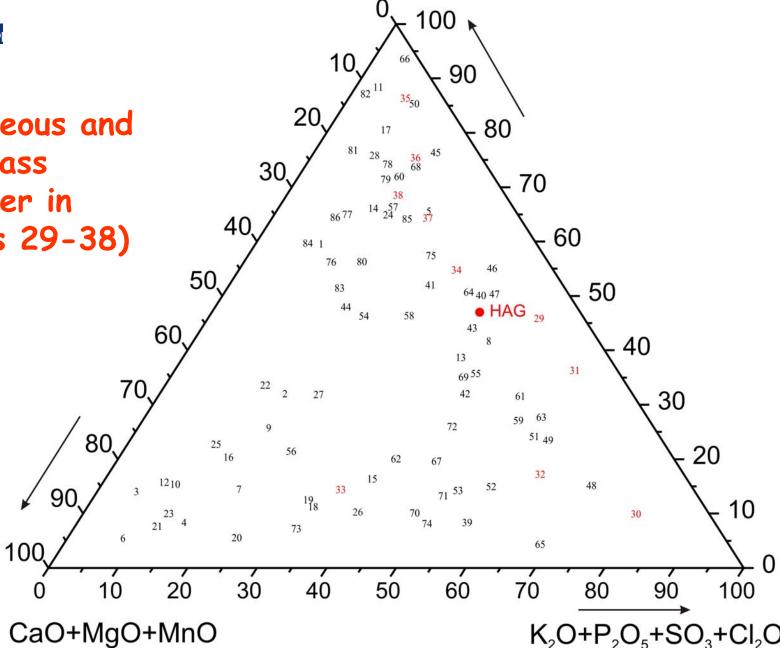
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SiO₂+Al₂O₃+Fe₂O₃+Na₂O+TiO₂

HAG - herbaceous and agricultural grass inorganic matter in ashes (samples 29-38)





SiO₂+Al₂O₃+Fe₂O₃+Na₂O+TiO₂

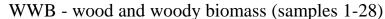
100

90

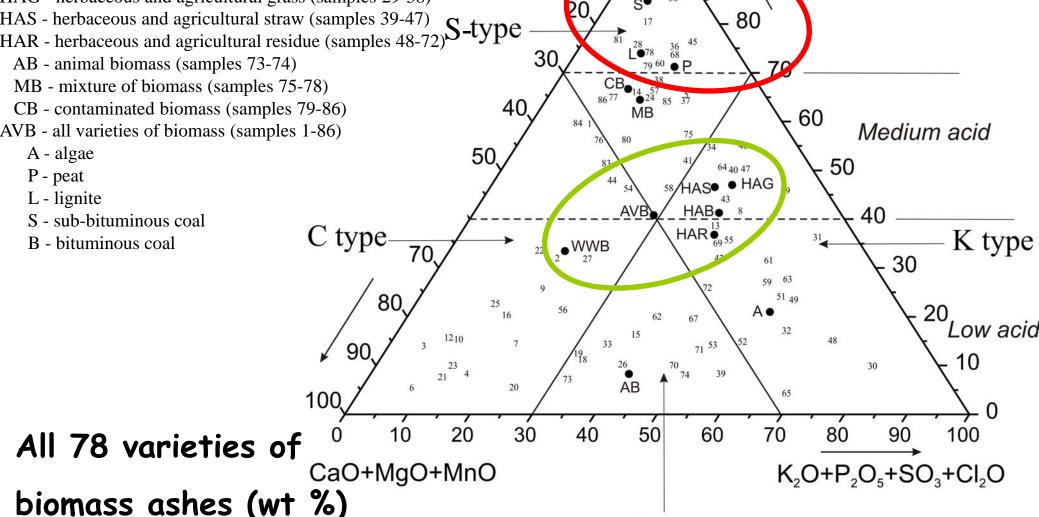
Nigh acid







- HAB herbaceous and agricultural biomass (samples 29-72)
- HAG herbaceous and agricultural grass (samples 29-38)
- HAS herbaceous and agricultural straw (samples 39-47)
- AB animal biomass (samples 73-74)
- MB mixture of biomass (samples 75-78)
- CB contaminated biomass (samples 79-86)
- AVB all varieties of biomass (samples 1-86)
 - A algae
 - P peat
 - L lignite
 - S sub-bituminous coal
 - B bituminous coal



CK type

All 78 varieties of

biomass ashes (wt



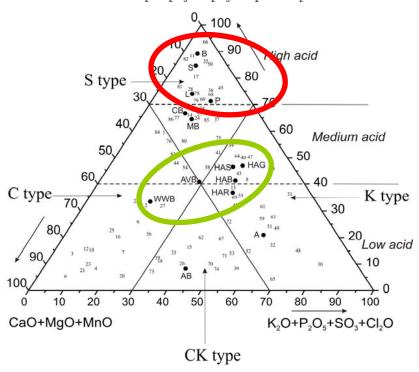
Preliminary Classification



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Preliminary chemical classification system for the inorganic matter of biomass:

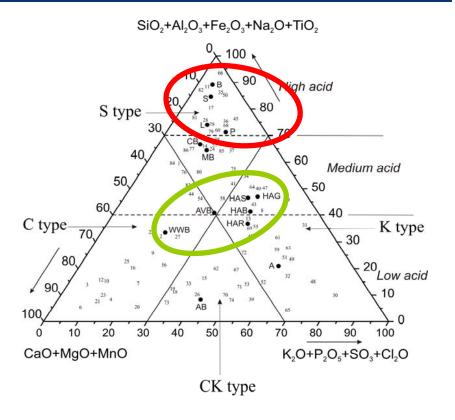
- (1) all varieties of biomass (close to the centre of the triangular graph);
- (2) WWB (Ca + Mg + Mn oxides above 30%);
- (3) HAB (K + P + S + Cl oxides above 30%);
- (4) SFFs (Si + Al + Fe + Na + Ti oxides above 70%);
- (5) **CB** (Si + Al + Fe + Na + Ti oxides between 40 and 70%)



Observations



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- (1) closer position of peat to coals than biomass (in contrast to proximate and ultimate composition)
- (2) differentiations among the biomass groups and some similarities for HAB sub-groups (HAG and HAS could be combined into one sub-group)
- (3) possibility for identification of some contamination in biomass fuels (unusual position of some samples 1, 5, 8, 11, 13, 14, 17, 24, 28, 57, 68)
- (4) position of marine macroalgae in "K ash type" with low acid tendency and highly enriched in Cl, S, Na, and Mg in comparison with the terrestrial biomass



Conclusions & Future Work



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The bulk chemical composition of biomass provides little information about the properties of biomass

Significant knowledge of the phase-mineral composition of biomass is essential in determining properties (analysis of biomass dry and dry ash-free bases essential)

Correlations exist between compounds and these correlations can be used to build a classification system (classification used for comparison with solid fossil fuels - for co-firing applications - application of ashes)

Future work will extend database for biomass and biomass ash for characterization of their:

- (1) phase-mineral composition and modes of element occurrence
- (2) thermal behaviour of phases and minerals
- (3) trace element contents and occurrences
- (4) traditional uses of ashes and consideration of innovative uses
- (5) potential advanced and environmentally safe applications





Thank you for your attention



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HAR - herbaceous and agricultural residue (samples 48-72)

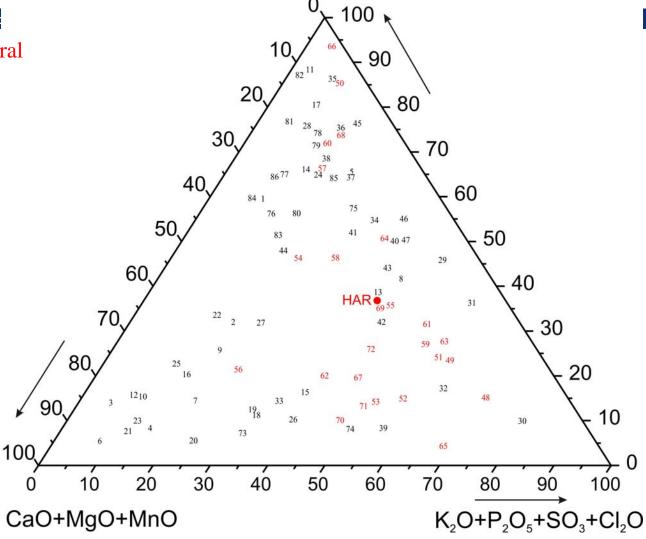


Fig. 4. Chemical classification system of the inorganic matter in biomass ashes based on 78 varieties of biomass (wt %)





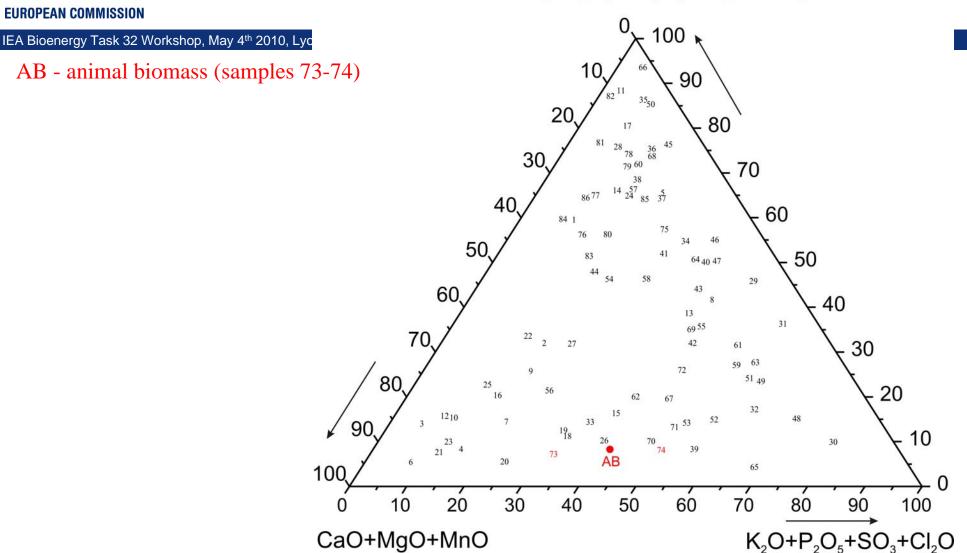


Fig. 4. Chemical classification system of the inorganic matter in biomass ashes based on 78 varieties of biomass (wt %)



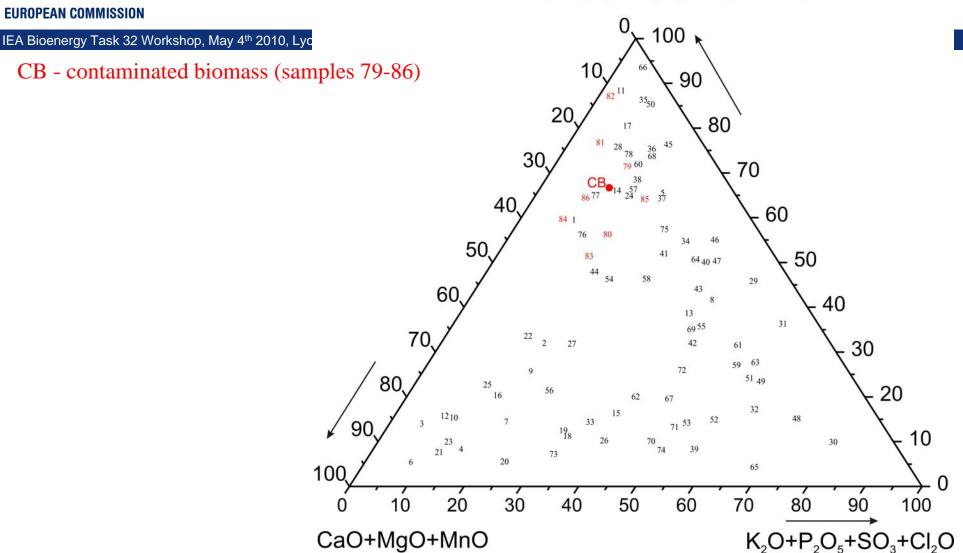


Fig. 4. Chemical classification system of the inorganic matter in biomass ashes based on 78 varieties of biomass (wt %)

