A novel biorefinery concept based on marginally used halophyte biomass

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Figure S1: Base peak chromatogram of S. ramosissima pretreated at 180 °C for 30 min with 60% ethanol; the MS2 inserts show the major glycans at m/z 367 and 353. Optional interpretations are shown.



Figure S2: Combined mass spectra and base peak chromatogram of S. ramosissima pretreated at 160 °C for 30 min with 60% ethanol; the MS2 inserts show the major glycans at m/z 501 and 563. Optional interpretations are shown.



Figure S3: Base peak chromatograms of S. ramosissima pretreated at 180 °C for 15 min with 40% and 60% ethanol, as well as at 180 °C for 30 min with 40% ethanol. The samples were analyzed in negative ion mode.



Figure S4: HSQC and quantitative ¹³C NMR spectra of lignin sample 0A4.



Figure S5: HSQC and quantitative ¹³C NMR spectra of lignin sample 0A6.



Figure S6: HSQC and quantitative ¹³C NMR spectra of lignin sample 0B4.



Figure S7: HSQC and quantitative ¹³C NMR spectra of lignin sample OB6.



Figure S8: HSQC and quantitative ¹³C NMR spectra of lignin sample 0C4.



Figure S9: HSQC and quantitative ¹³C NMR spectra of lignin sample 0C6.



Figure S10: HSQC and quantitative ¹³C NMR spectra of lignin sample 1A4.



Figure S11: HSQC and quantitative ¹³C NMR spectra of lignin sample 1A6.



Figure S12: HSQC and quantitative ¹³C NMR spectra of lignin sample 1B4.



Figure S13: HSQC and quantitative ¹³C NMR spectra of lignin sample 1B6.



Figure S14: HSQC and quantitative ¹³C NMR spectra of lignin sample 1C4.



Figure S15: HSQC and quantitative ¹³C NMR spectra of lignin sample 1C6.



Figure S16: HSQC and quantitative ¹³C NMR spectra of lignin sample 1D4.





Figure S17: HSQC and quantitative ¹³C NMR spectra of lignin sample 1D6.



Figure S18: HSQC and quantitative ¹³C NMR spectra of lignin sample 2B4.



Figure S19: HSQC and quantitative ¹³C NMR spectra of lignin sample 2B6.