

Peculiarities of Synthesis and Properties of Lignin-Silica Nanocomposites Prepared by Sol-Gel Method

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Received: date; Accepted: date; Published: date

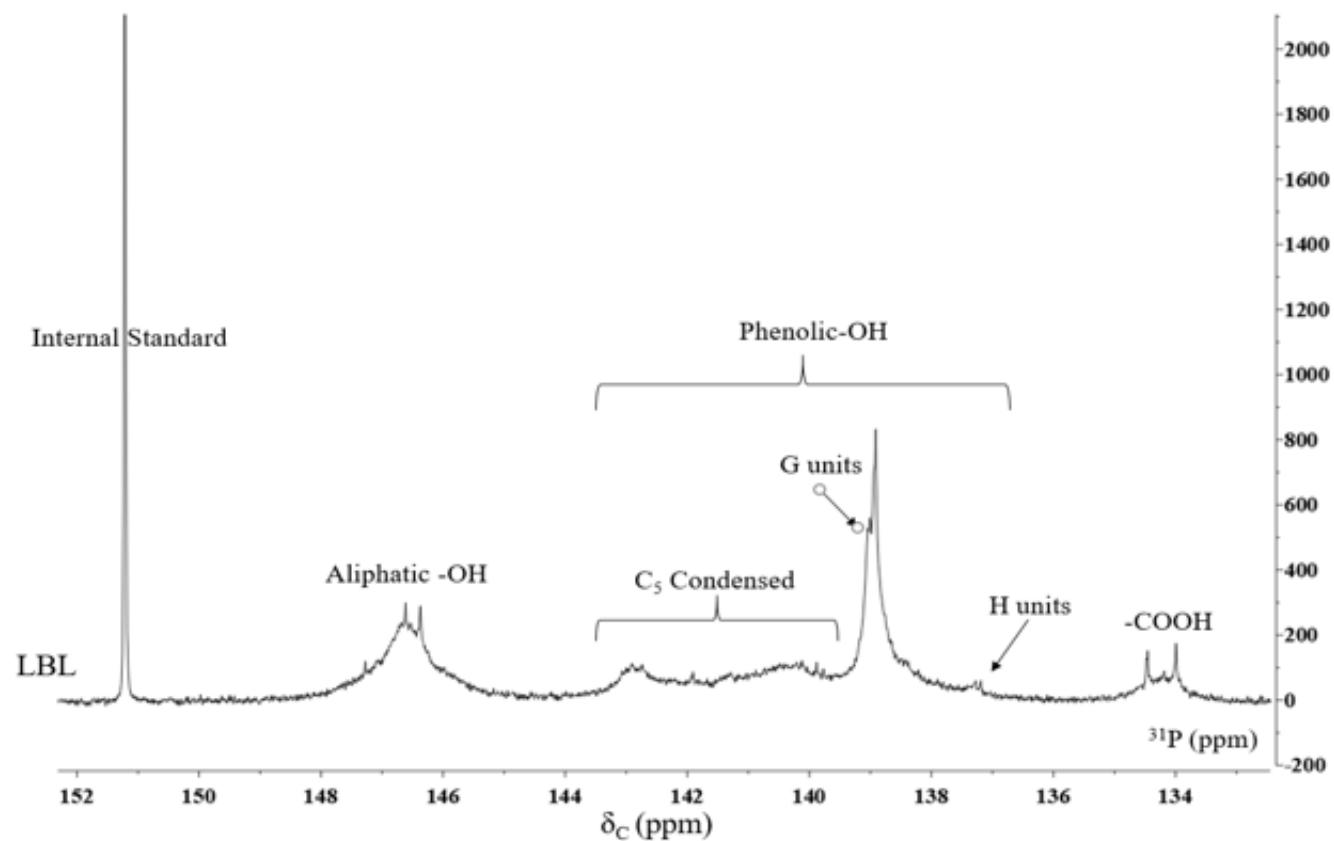
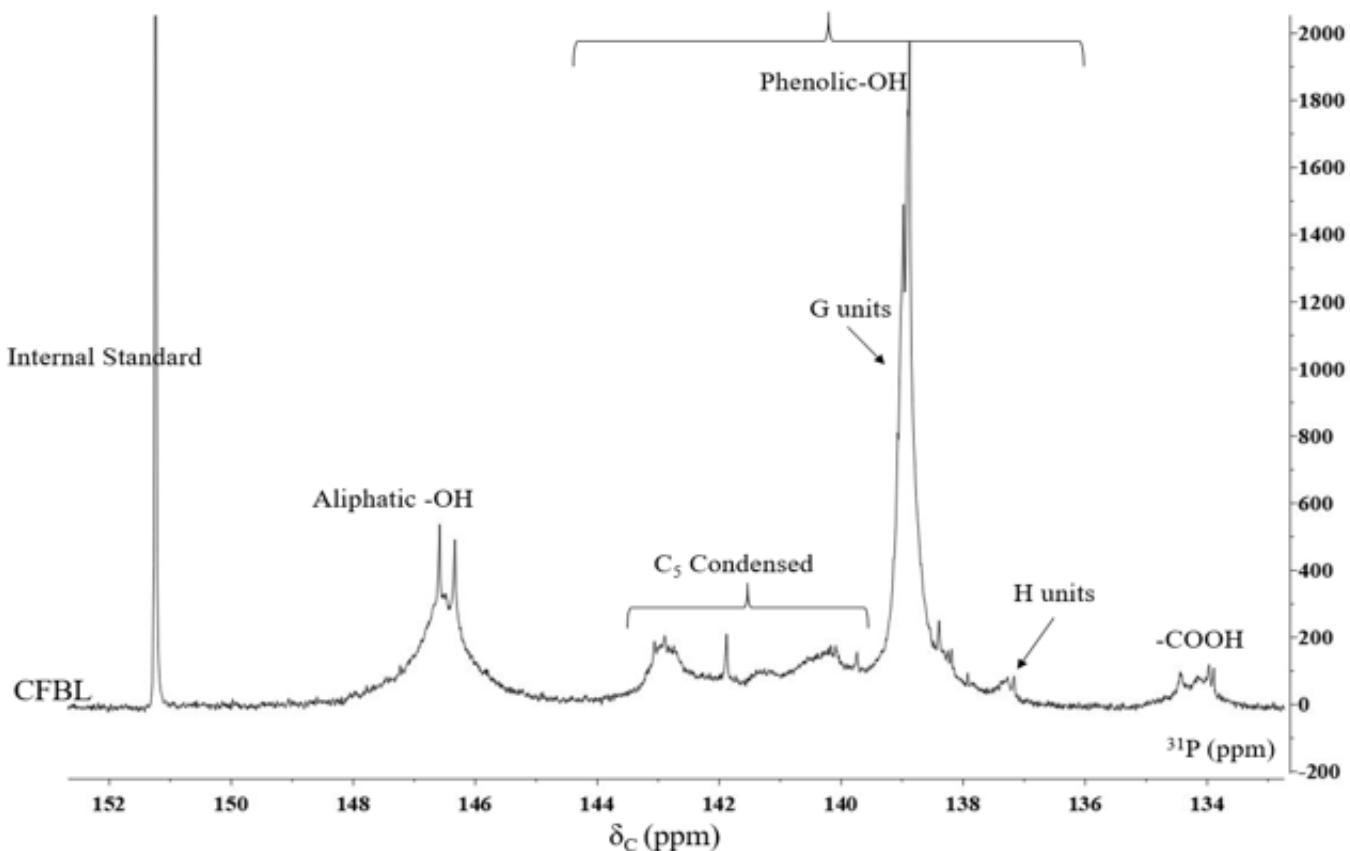
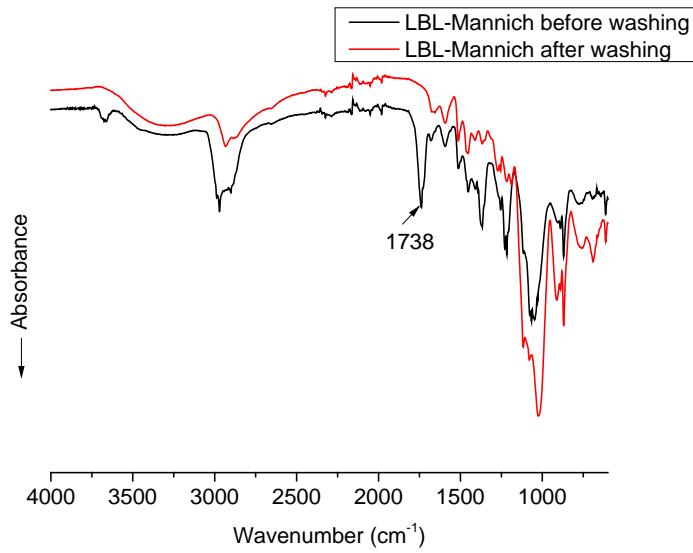
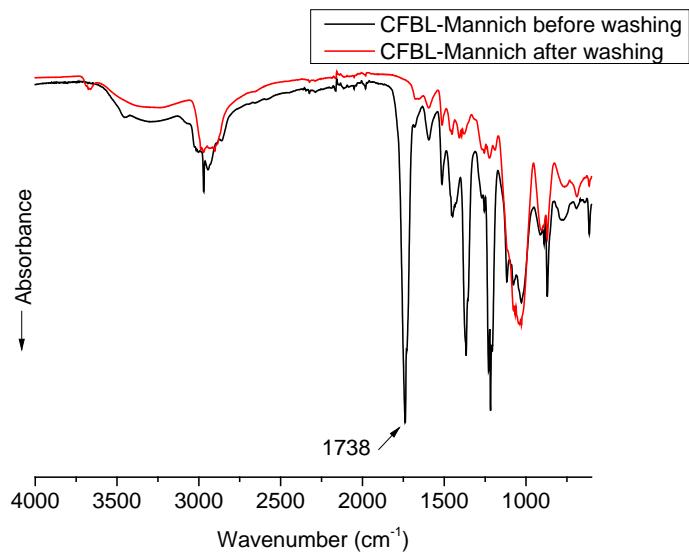


Figure S1. ^{31}P -NMR spectra of the CFBL and LBL.

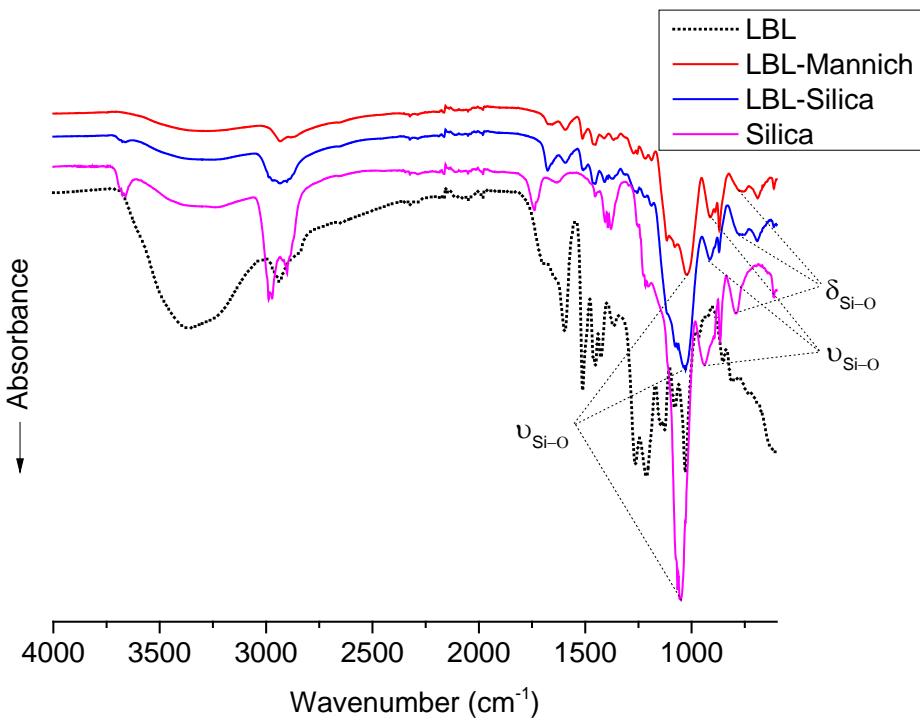


a

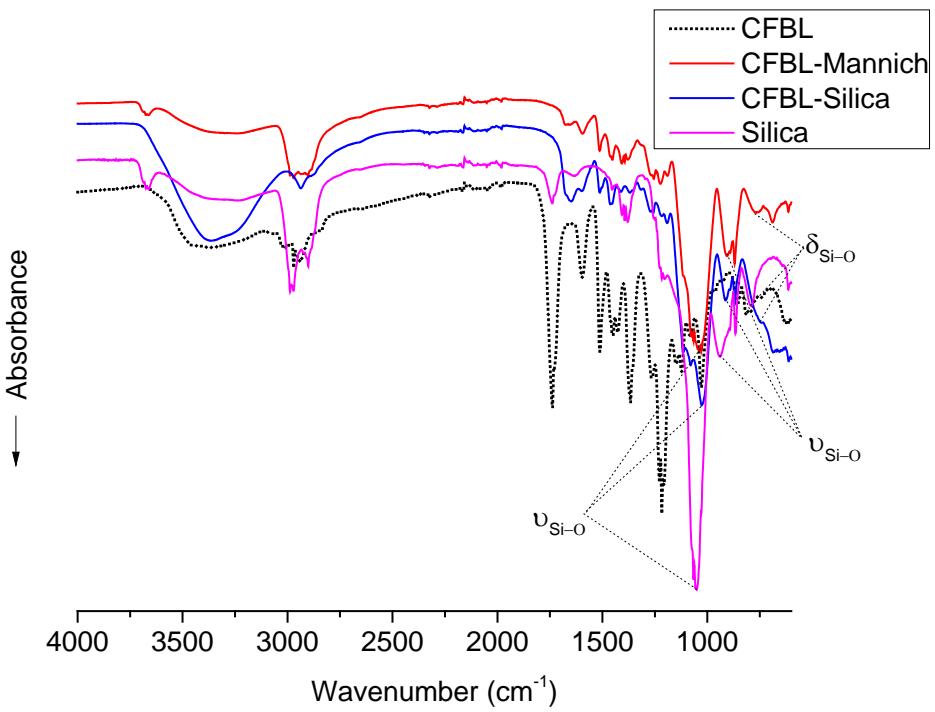


b

Figure S2. FTIR spectra of the LBL-Mannich (a) and CFBL-Mannich (b) samples before and after precipitation and washing by diethyl ether.

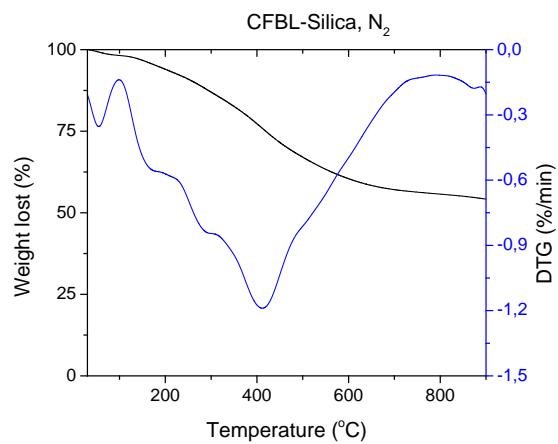
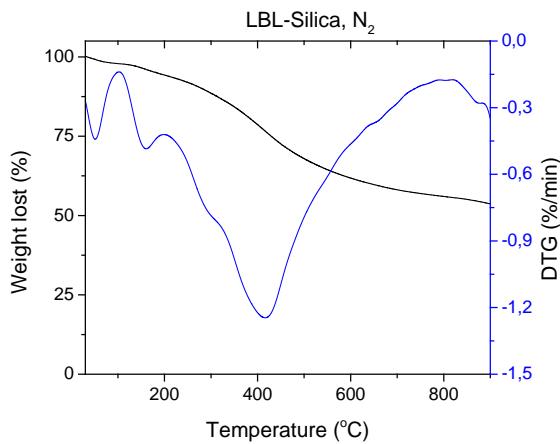
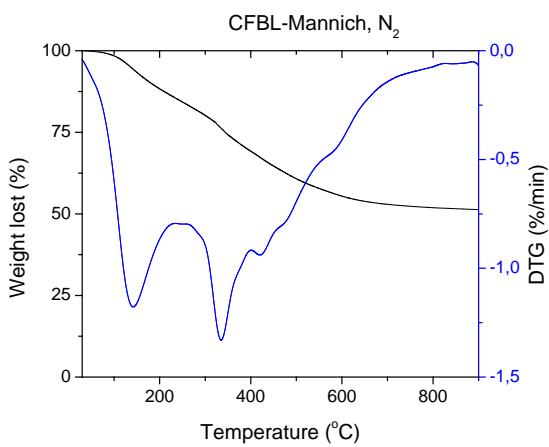
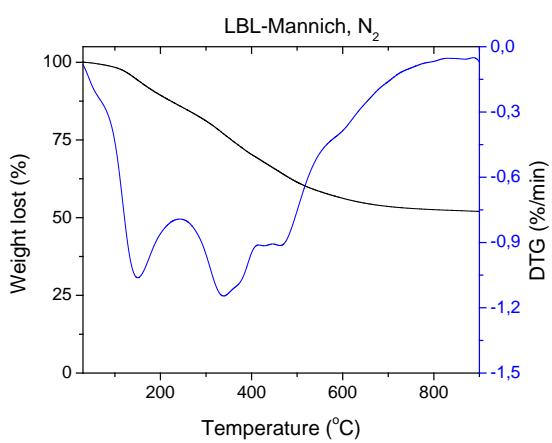
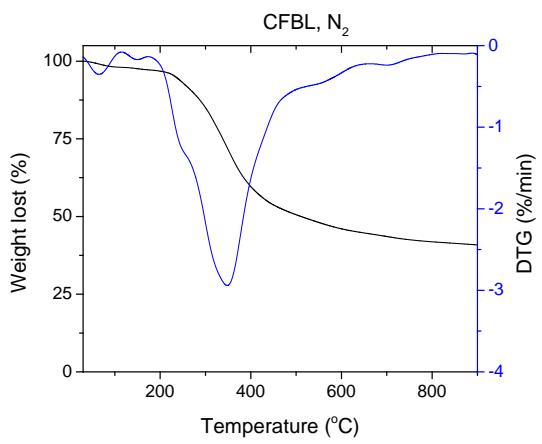
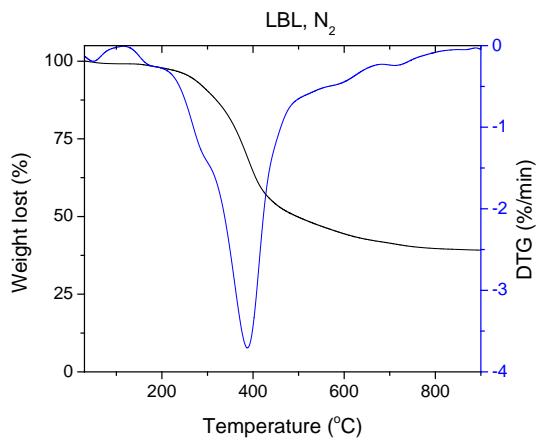


a



b

Figure S3. FTIR spectra (in range 4000–600 cm⁻¹) of the materials based on LBL (**a**) and CFBL (**b**): initial lignin; lignin modified by the Mannich reaction; lignin–silica hybrids and synthesized pure silica.



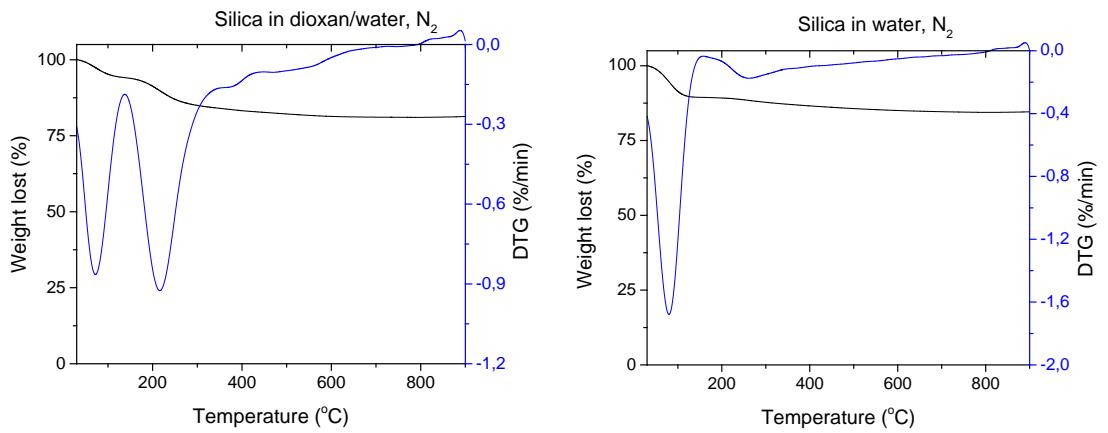
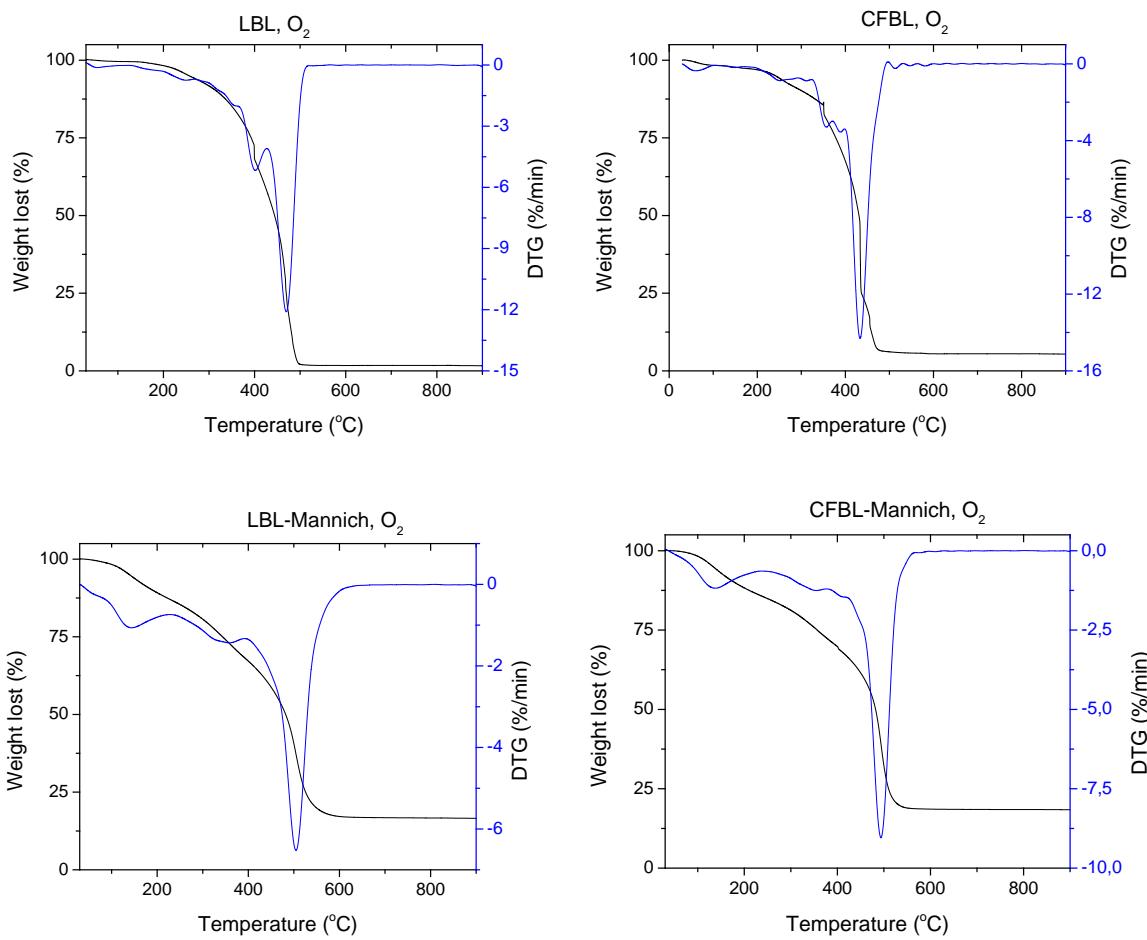


Figure S4. TG and DTG curves of thermal decomposition in N₂ atmosphere for initial and modified lignins, hybrid composites, and silica.



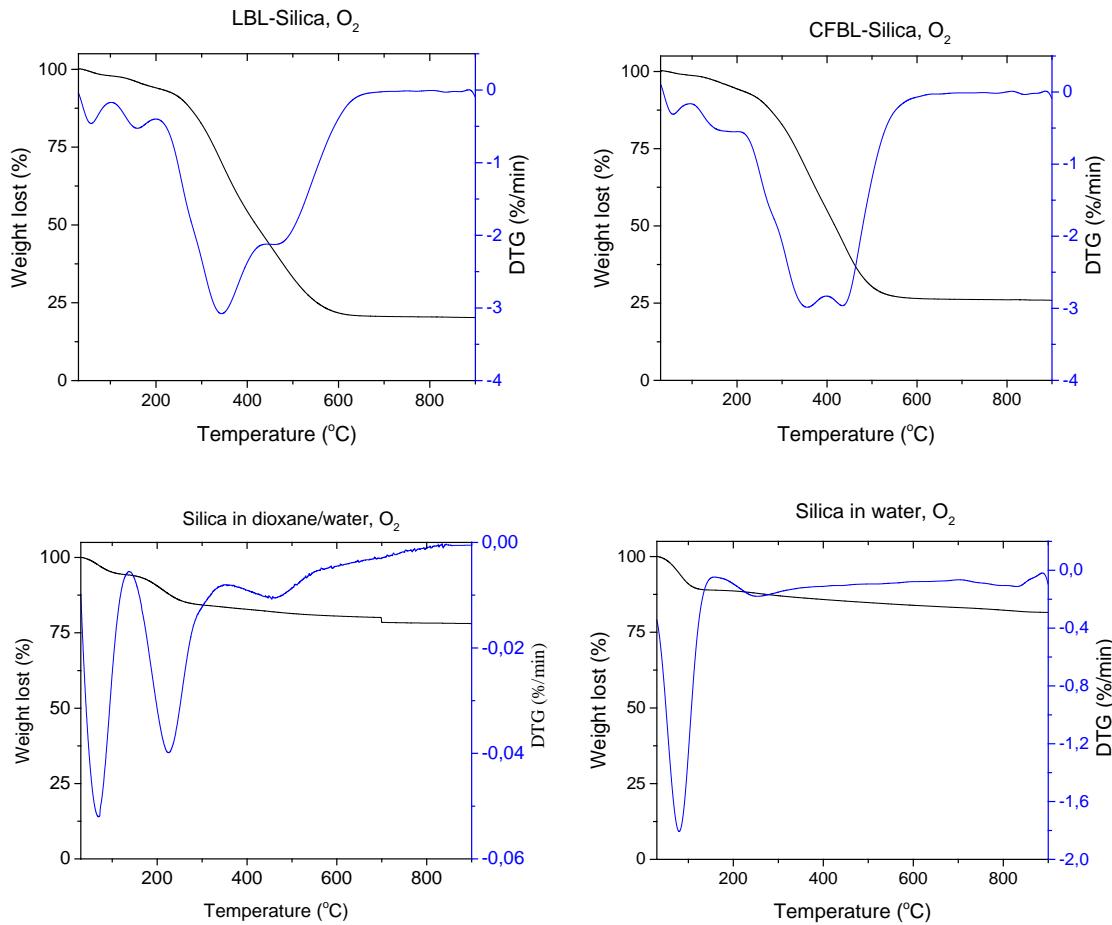


Figure S5. TG and DTG curves of thermal decomposition in O₂ atmosphere for initial and modified lignins, hybrids, and silica.

Table S1. Characteristics of thermal decomposition of studied materials in O₂ atmosphere.

| Material | T _{max} , °C (DTG) | Δm, % (TG) | Δm _{total} , % (TG) | Process |
|--------------|--------------------------------|------------|---------------------------------|----------------------------------------------|
| LBL | 56 | 0.3 | 97.77 | Moisture evaporation |
| | 250 | 6.4 | | Lignin decomposition onset |
| | 401 | 16.3 | | Lignin decomposition |
| | 468 | 61.6 | | Lignin combustion |
| CFBL | 58 | 0.7 | 94.5 | Moisture evaporation |
| | 247 | 6 | | Lignin decomposition onset |
| | 383 | 27 | | Lignin decomposition |
| | 435 | 63.4 | | Lignin combustion |
| LBL-Mannich | 54 | 0.5 | 83.4 | Moisture evaporation |
| | 142 | 5.2 | | Water and ethoxyl radical elimination |
| | 347 | 26.1 | | Decomposition of the aminopropyl radical |
| | 504 | 61.2 | | Lignin decomposition and combustion |
| CFBL-Mannich | 138 | 13.3 | 82.3 | Ethoxyl radical elimination |
| | 354 | 19.7 | | Decomposition of the aminopropyl radical |
| | 493 | 49.2 | | Lignin decomposition and combustion |
| LBL-Silica | 59 | 1.4 | 77 | Moisture evaporation |
| | 158 | 5.1 | | Water and ethoxyl radical elimination |
| | 344 | 45.9 | | Lignin and aminopropyl radical decomposition |
| | 470 | 29.4 | | Lignin combustion |
| CFBL-Silica | 58 | 1.4 | 73 | Moisture evaporation |
| | 158 | 5.1 | | Water and ethoxyl radical elimination |
| | 353 | 38.44 | | Lignin and aminopropyl radical decomposition |
| | 435 | 28.3 | | Lignin combustion |

| | | | | |
|----------------------------|-----------|--------------|------|--------------------------------------------------------------------------------------|
| Silica in water | 67 210 | 11 2 | 22.1 | Physically adsorbed water evaporation; condensation of the silica hydroxyl groups |
| Silica in dioxane/water | 80 248 | 5.4 10.75 | 18.3 | Dioxan evaporation; condensation of the silica hydroxyl groups |

Adsorption of methylene blue dye

The synthesized hybrid composites were tested as sorbents for the removal of methylene blue dye from aqueous solutions. This study was conducted to evaluate the dependence of the sorption properties of the synthesized hybrid materials on the structure of the lignin component and to compare the sorption properties of the initial LignoBoost and CleanFlowBlack lignins.

The adsorption capacity of the hybrids, initial lignins, and silica were estimated from the adsorption isotherms of methylene blue dye in a neutral medium. The equilibrium studies were conducted over 48 h for all systems. The obtained results were compared and are shown in a diagram in Figure 10. As seen from Figure 10, the adsorption capacity of the hybrid composites is higher than that of the original lignins. In the case of LBL (adsorption capacity = $31.2 \text{ mg}\cdot\text{g}^{-1}$), the increase reached 30% (adsorption capacity of LBL–silica = $41.6 \text{ mg}\cdot\text{g}^{-1}$), and in the case of CFBL (adsorption capacity of CFBL–silica = $32.3 \text{ mg}\cdot\text{g}^{-1}$), the adsorption capacity almost doubled (adsorption capacity = $58.9 \text{ mg}\cdot\text{g}^{-1}$). It could be concluded that the increase in the adsorption capacity is presumably related to an increase in the organophilic properties of the hybrid composites. The highest sorption of the studied dye was observed for the CFBL–silica composite. The lignin in this composite has a less condensed structure, a lower molecular weight, and a larger number of functional groups. Based on the higher reactivity, the lignin was probably more homogeneously distributed on the surface of the hybrid composite, which resulted in better accessibility of the functional groups that participated in the sorption process.

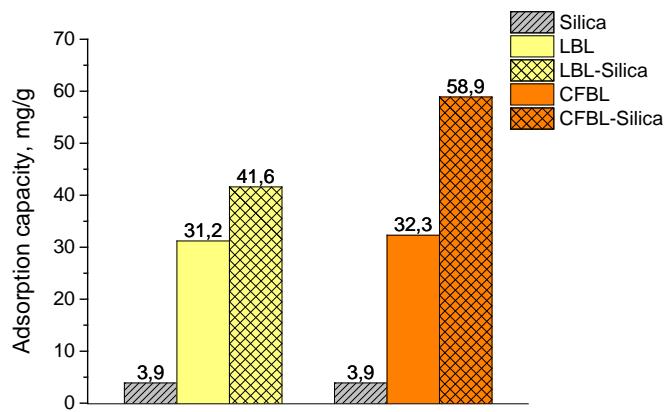


Figure S6. Adsorption capacities of the synthesized silica, initial CFBL and LBL, and LBL–silica and CFBL–silica composites for methylene blue dye in a neutral medium.