

Biomass Pyrolysis in a Heated Microtubular Reactor

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A heated SiC microtubular reactor has been developed to decompose biomass monomers such as CH₃CHO, C₆H₅OH, C₆H₅OCH₃, HOC₆H₄OCH₃, C₆H₅CH₂CH₂OC₆H₅, and furan. The pyrolysis experiments are carried out by passing a dilute mixture of the organic substrate (roughly 0.1 — 1 %) entrained in a stream of a buffer gas (either He or Ar) through a heated SiC reactor that is 2-3 cm long and 1 mm in diam. Common pressures in the reactor are 50 – 200 Torr and the SiC tube is heated in the range of 1200 — 1900 K. Typical transit times through the reactor are 50 — 200 μsec after which the gas mixture emerges as a skimmed molecular beam at a pressure of approximately 10 μTorr and all chemistry is quenched. The reactor has been deployed in pulsed and CW configurations. In Colorado a pulsed reactor is used because the pyrolysis products are identified by photoionization mass spectroscopy with a 10 Hz YAG laser at λ₀ = 118.2 nm or 10.487 eV. Separate experiments use matrix infrared absorption spectroscopy to identify the pyrolysis products and to confirm the assignments of the PIMS. In Calif. a CW reactor is used because a CW synchrotron is used as the light source for the PIMS. The pyrolysis of CH₃CHO will be discussed. We observe CH₃CHO (+M) → CH₃ + H + CO + CH₂=C=O, CH₂=CHOH, HC≡CH + H₂O.

μTubular reactor cracking:
CH₃CHO + Δ → products

4 Torr CH₃CHO (0.3 %) entrained in
2 atm He carrier gas behind pulsed
valve (open time 1 msec @ 10 Hz)

1 mm x 2 cm SiC tube @ 300–1700 K

IE(CH₃CHO) = 10.2298 ± 0.0007 eV

ħω_{VUV} + CH₃CHO → CH₃CHO⁺ + e⁻

118.2 nm VUV Laser (10.487 eV) @ 40 cm downstream

CH₃CHO⁺
TOF detection

Supersonic Jet
Radicals/He into
10⁻⁷ Torr

T
H
U
R
S
D
A
Y