

Influence of Supercritical CO₂ and N₂ on the Viscosity and Solidification Behavior of Polyethylene

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The viscosity of polymer melts plays a critical role in processing efficiency, with high viscosities requiring increased energy input. Supercritical carbon dioxide (scCO₂) offers a promising approach to reduce melt viscosity without the need for solvent-based plasticizers [1]. In this study, the rheological behavior of different polyethylene (PE) types was investigated under elevated pressure using a plate-plate pressure cell capable of both rotational and oscillatory measurements.

Flow curves, frequency sweeps and temperature ramps were performed at ambient pressure, 120 bar N₂, and 120 bar scCO₂. While N₂ showed no significant impact on melt viscosity, it increased the solidification temperature, depending on the PE type. In contrast, scCO₂ acted as a temporary plasticizer, substantially lowering melt viscosity and slightly delaying crystallization. Oscillatory tests proved particularly useful for high-viscosity samples, enabling reliable measurements even near the melting point, provided strain and frequency were carefully adjusted.

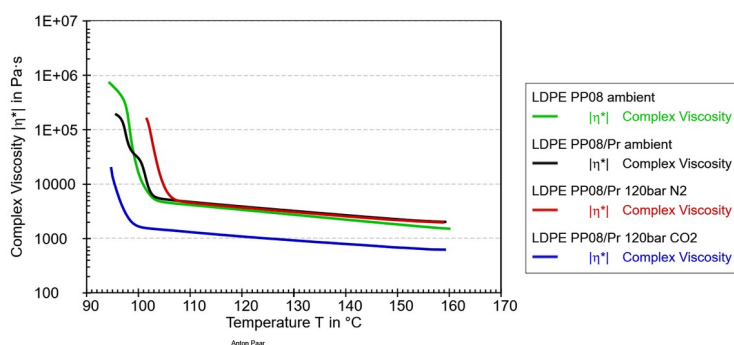


Figure 1: Temperature ramp in oscillation of LDPE at ambient conditions, 120 bar N₂ and 120 bar CO₂

The results highlight the potential of scCO₂ in energy-efficient polymer processing and foaming applications, and demonstrate the value of high-pressure rheometry for characterizing pressure-dependent polymer behavior.

Keywords: Polyethylene Viscosity, High-Pressure Rheology, Supercritical CO₂, Polymer Crystallization, Plasticization with CO₂

References

[1] Wan, C., Lu, Y., Liu, T., Zhao, L., & Yuan, W. Foaming of low density polyethylene with carbon dioxide based on its in situ crystallization behavior characterized by high-pressure rheometer. *Industrial & Engineering Chemistry Research* **2017**, 61(16), 5334-5345.