

Synthesis and Characterization of the Terpolymer of Itaconic Acid with Acrylamide and 2-Acrylamido-2-methylpropanesulfonic Acid

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Abstract: The terpolymer of itaconic acid, acrylamide and 2-acrylamido-2-methyl-1-propane sulfonic acid was synthesized through the free-radical polymerization. The IR spectra confirmed that there was no olefinic band, while the TGA results revealed that the terpolymer was of high thermal stability.

Keywords: Itaconic acid, acrylamide, 2-acrylamido-2-methyl-1-propane sulfonic acid, TGA.

The thermal stability of the polymer is of great significance in paper, ceramics and oil industry¹. Conventional polymers such as polyanionic cellulose can not meet high temperature limitations^{2,4}. Aggour⁵⁻⁶ and Collette⁷ *etc.* have conducted extensive research on synthesis and characterizations of the copolymer or terpolymer containing 2-acrylamido-2-methyl-1-propane sulfonic acid (AMPS). Till now, very few research works have been reported on the terpolymer of itaconic acid (IA), acrylamide (AM) and AMPS.

The polymerization of the terpolymer was carried out under the nitrogen atmosphere. 425 g of deionized water, 200 g of a 50% solution of AM, 30 g of a 40% solution of disodium IA and 276 g of a 50% solution of AMPS were added into a 1 L reactor. Then 52.5 g of a 5% solution of sodium persulfate was drop-fed into the reactor uniformly over a period of 2 h at 75°C. At the end of the initiator addition, 75°C was maintained for an additional 2 h. The precipitated polymer was isolated by 2 L acetone and dried under vacuum at 40°C. The molecular weight of the terpolymer was about 500,000 as measured by the limiting viscosity method and calculated through the relation as follows:

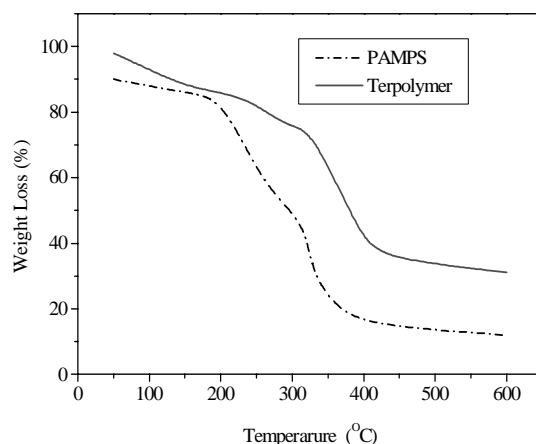
$$[\eta] = 3.73 \times 10^{-4} M_w^{0.66}$$

In the IR spectra of the IA-AM-AMPS terpolymer, there was a band at 1406 cm⁻¹ due to C-O stretching of COO⁻ groups. The AMPS units were revealed by CO at 1558 cm⁻¹, SO₂ at 1217 cm⁻¹, CH at 1451 cm⁻¹ and NH at 3211 cm⁻¹. The AM units were characterized by CO at 1665 cm⁻¹, CH at 579 cm⁻¹ and NH₂ at 3339 cm⁻¹. There was no olefinic band at 1635-1620 cm⁻¹.

The TGA curves for the IA-AM-AMPS terpolymer and the AMPS homopolymer

(PAMPS) were showed in **Figure 1**. Obviously, the terpolymer was of higher thermal stability than PAMPS. The thermal degradation of the terpolymer became appreciable at approximately 240°C while PAMPS began to degrade at 182°C. The terpolymer sample turned yellow at first, later fused, and the rate of degradation gradually decreased. The terpolymer exhibited three degradation stages. The first degradation stage occurred in the range of 240~330°C corresponding to the decomposition of amide and carboxyl groups. The second stage appeared at 340~420°C due to the decomposition of sulfonic groups, while the third stage demonstrated smoothly at the higher temperature than 430°C in correspondence with the decomposition of the terpolymer backbone.

Figure 1 TGA curves for the IA-AM-AMPS terpolymer and PAMPS



Acknowledgment

We are grateful for the financial support of the Shengli Oilfield.

References

1. L. L. Carney, *J. Pet. Tech.*, **1980**, 32, 385.
2. J. Chatterji, J. K. Borchardt, *J. Pet. Tech.*, **1981**, 33, 2042.
3. H. Kheradmand, J. Francois, V. Plazanet, *Polymer*, **1988**, 29, 860.
4. W. O. Parker, A. Lezzi, *Polymer*, **1993**, 34, 4913.
5. Y. A. Aggour, *Polymer Degradation and Stability*, **1994**, 45, 273.
6. Y. A. Aggour, *Polymer Degradation and Stability*, **1994**, 44, 71.
7. C. Collette, F. Lafuma, R. Audibert, R. Brouard, *J Appl Polym Sci*, **1994**, 53, 755.

Received 20 October, 2000