Graft Copolymerization of Acrylonitrile and Ethyl Methacrylate on Dextrin

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ABSTRACT

Graft copolymers of Acrylonitrile and ethyl methacrylate on dextrin were prepared by the use of ceric ion initiator in aqueous medium at 29°C. The molecular weight of grafted poly(ethyl methacrylate) chains were higher than for polyacrylonitrile grafts; but the latter were more frequently grafted on the backbone polymer. The dependence of the molecular weight of the grafted chains on ceric ion concentration suggests that interaction of growing graft polymer chains with ceric ion is important in the termination process of graft copolymerisation. Graft copolymerisation of ethyl methacrylate onto dextrin in the presence of various amounts of acrylonitrile was studied. The compositions of the copolymer samples, calculated from their nitrogen content, showed that in the presence of ethyl acrylate, relatively small amounts of acrylonitrile monomer was grafted onto dextrin.

INTRODUCTION

Ceric ion has been used to initiate graft copolymerisation of many monomers on various substrates1–7. Grafting is generally considered to result from propagation by radicals generated on the backbone polymer. It is well accepted that the extent of graft polymer formation depends on the nature and reactivity of the backbone polymer towards initiation of graft polymer formation, the reactivity of the active sites towards propagation, the nature of termination reactions of copolymerisation, and the polarity of the monomer. Several workers have reported on graft copolymerisation onto starch8–10. Graft copolymers of polysaccharides are of interest11. This communication examines some aspects of graft copolymerisation onto dextrin.

EXPERIMENTAL

Acrylonitrile (AN) and ethyl methacrylate (EMA) monomers were purified from hydroquinone stabiliser by extraction with aqueous alkaline solution followed by distillation. Ceric ammonium nitrate was used without further purification. Dextrin was used without further purification. Dextrin was prepared from cassava starch using 20% (v/v) hydrochloric acid12. Cassava starch (20.0g) was moistened with 5ml of dilute aqueous hydrochloric acid (0.2 vol. %), spread on a tray and heated at 60°C for 2hr in an oven. The acidified starch was further heated at 100°C for 5hrs and 100°C for another 5hrs; and cooled in a desiccator. Graft copolymerisation was based on the method described by Lepoutre and Hui13 for grafting onto woodpulp.

A known amount of dextrin (0.5g) was dissolved in 100ml of deionised water containing a definite amount of ceric ion. The initiator was allowed to interact with dextrin for 30 mins and then a definite amount of the monomer was added dropwise. The polymerisation was allowed to proceed for 4hrs and then stopped by the addition of 5ml of 2% (v/v) wuinol solution to the reaction mixture. The reaction mixture was poured into an excess of methanol and filtered. The residue was air-dried and weighed. The ungrafted poly(ethyl
methacrylate) (PEMA) and polyacrylonitrile (PAN) homopolymers were extracted with benzene and dimethyl formamide respectively. The dextrin graft copolymer was air-dried and weighed. The percentage graft level, (P_g), was calculated from the weights of dextrin and grafting products.

\[
P_g = \frac{\text{weight of grafted polymer} \times 100}{\text{Weight of dextrin}}
\]

Graft copolymerization of ethylacrylate (0.24m) in the presence of various amounts of acrylonitrile using 0.2M ceric ion was also studied at 29°C. The grafted PEMA and PAN chains were isolated from the backbone polymer by treatment with hydrochloric acid. The isolated PEMA and PAN grafts were purified by solution in benzene and dimethylformamide respectively and precipitation with methanol and airdried. The molecular weights of the polymers were determined from viscosity measurement using the following relationships

\[
\log(\eta) = \log 4.75 \times 10^{-4} + 0.50M_v \quad (2)
\]
\[
\log(\eta) = \log 3.35 \times 10^{-4} + 0.72M_v \quad (3)
\]

for PEMA in isopropanol at 30°C and PAN in dimethylformamide at 30°C respectively. The frequency of grafting (F_g), defined as the number of polymer grafts per 10^4 anhydroglucose units (AGU), was calculated using the relationships

\[
F_g = \frac{(\text{weight of grafted polymer})(\text{mw of AGU}) \times 10^4}{\text{mw of grafted polymer}} \quad \ldots (4)
\]

The nitrogen content of the grafted PEMA-PAN copolymer samples was determined by the Kjedahl method. The copolymer compositions were worked out from the nitrogen content of the samples using the relationship

\[
\text{mole fraction of AN in copolymer} = \frac{(BxN)/14B^{-1}}{1 - (BxN)/14C^{-1} + (BxN)/14B^{-1}} \quad \ldots (5)
\]

Here N is the weight of nitrogen per gram of copolymer sample, B is the molecular weight of acrylonitrile and C is the molecular weight of ethyl methacrylate.

**RESULTS AND DISCUSSION**

The variation of P_g with ceric ion concentration using 0.16M EMA and 0.46M AN is shown in Fig. 1. The results show that the initial increase in P_g with ceric ion concentration attained maximum values of about 11% and 225% for the graft copolymerisation of EMA and AN respectively at 0.2M ceric ion and decrease with further increase in ceric ion concentration. The low values of P_g at high initiator concentration have been associated with the formation of ceric ion complex with the backbone polymer which reduces both the number of grafting sites on the backbone polymer and the amount of ceric ion available to initiate graft polymer formation. The values of P_g obtained for dextrin are much higher than the values reported for starch.
Fig 1: Grafting of EMA (o) and AN (î) on dextrin at 29°C using 0.16M EMA and 0.46M AN

The molecular weights of grafted PEMA and PAN chains at various ceric ion concentrations are shown in Fig. 2. The results show that the molecular weight of grafted PEMA chains increased with increase in initiator concentration up to 0.2M ceric ion and decreased with further increase in ceric ion concentration. The molecular weights of grafted PAN chains, on the other hand, decreased with increase in ceric ion concentration. The dependence of the molecular weight of the grafted chains on ceric ion concentration suggests that termination of graft copolymerisation by the interaction of growing graft polymer chains with ceric ion would be the main termination process in the copolymerisation of AN and would become significant at relatively high ceric ion concentration. It is well recognized that in such polymerisation, where mutual termination of growing polymer chains are low, the chain transfer constant (Kₐ/Kₚ), can be determined from the plot of 1/N against ceric ion concentration using the relationship

\[ \frac{1}{N} = \frac{1}{N_0} + K_A \frac{[Ce^{4+}]}{K_p[M]} \]  

Here N and N₀ are the degrees of graft copolymerisation in the presence and absence of ceric ion respectively.

It can be seen from the plots in Fig. 3 that termination of copolymerisation chain transfer to ceric ion is important at high initiator concentrations. The values of the transfer constant were determined to be 3.68 x 10⁻⁵ and 3.30 x 10⁻⁴ for the graft copolymerisation of EMA and AN respectively.

Fig 2: Dependence of the Molecular weight grafted PEMA (o) and PAN (î) chains on ceric ion concentration

Fig 3: Chain transfer to ceric ion in the graft copolymerization at EMA (o) and AN (î) on dextrin at 29°C
Li and Fu\textsuperscript{16} in their study of the graft copolymerisation of butyl acrylate on gelatin proposed relationships between the rate of graft copolymerisation (RG) and ceric ion concentration for cases where termination occurs by the interaction of two growing graft polymer chains as

\[
R_g = \frac{(k_p k_d / k_t_1)}{k_d k[Dex]^{1/2} + k_1[M]^{1/2}}
\] (7)

and where termination occurs by the interaction of ceric ion with growing graft polymer chains.

\[
R_g = \frac{1}{k_t_2} \left\{ M \right\}^2 \frac{[Dex]}{[M] + \left\{ k_0 / k_1 \right\} [Ce_4^+]}
\] (8)

Equations 7 and 8 predict that the kinetic order of the rate of graft polymerisation in initiator would be 0.5 and –1 for termination of copolymerisation brought about by mutual combination of growing polymer radicals and termination by chain transfer process respectively. The rate of graft copolymerisation can be obtained from the relationship.

\[
R_g = \frac{\text{weight of grafted polymer} \times 1000}{\text{(mw of monomer)(reaction time)(reaction volume)}}
\]

The effect of ceric ion concentration on the rate of graft copolymerisation of EMA and AN on dextrin is shown in Fig. 4. It can be seen that the plots of \( R_g \) against initiator concentration are linear at low ceric ion concentration, which agrees with mutual termination of growing polymer chains. At higher initiator concentration, the plots deviate from linearity, suggesting that termination of copolymerisation probably is due to transfer to ceric ion. It has been argued that for polymerisation in aqueous dextrin solution, the rates of mutual termination of growing graft polymer chains would be low and diffusion controlled. Such would be the case at low conversions of monomers to polymer and termination of graft copolymerization would proceed mainly by transfer reactions (9). Fig 5 shows the variation of \( F_g \) with ceric ion concentration. The number of grafted polymer molecules per \( 10^4 \), AGU, varies from about two to eleven and from

Fig 4: Dependence of rate of graft copolymerization of EMA (\( \circ \)) and AN (\( \bullet \)) on dextrin at 29\(^\circ\)C

Fig 5: Graft copolymerization onto dextrin Dependence of \( F_g \) of PEMA(\( \circ \)) and PAN (\( \bullet \)) on ceric ion concentration.
twenty-six to more than nine hundred for the graft copolymerisation of EMA and AN respectively on dextrin. Fanta and Coworkers\(^5\) in a study of copolymerisation of acrylonitrile on a variety of starch samples reported that graft copolymerisation in dilute aqueous starch dispersion enhanced the frequency of grafting. It is thought that the reactive end groups produced during the dextrinisation process would explain the fairly high frequencies of grafting on dextrin.

The influence of acrylonitrile on the grafting of EMA on dextrin is shown in Fig. 6. The results show that \(P_g\) increases with the AN content of the feed composition. The increase in \(P_g\) with increasing proportion of AN in the monomer mixture suggests that the reactivity of a growing graft copolymer chain is either accentuated by the presence of monomeric units of AN or that AN molecules are grafted independently of the EMA molecules.

The compositions of the copolymer samples were worked out from their nitrogen contents according to Eq. 5. The results show a general increase in the contents of monomer units of AN in the copolymer with increasing proportion of AN in the monomer mixture. However, it can be seen that the increase in AN monomeric units in the copolymer does not correspond to the increase of AN in the feed. For instance an increase in the amount of AN in the feed by as much as 40mol% resulted in less than 3mol% increase in the AN content of the copolymer. Acrylonitrile monomeric units can either be built into the graft copolymer chains randomly with EMA monomeric units or into PAN homopolymer chains. Either way, our method of analysis should account reasonably well for AN monomeric units in the grafting products. For example, determination of AN content of PAN homopolymer by analysis for nitrogen gave about 94\% accuracy\(^17\).

The low values of AN content of the copolymer samples even at fairly high AN levels in the feed are indicative of the low reactivity of acrylonitrile in copolymerisation reactions with acrylates\(^18,19\).

**REFERENCES**


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