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E. I. DU PONT DE NEMOURS & COMPANY

AIKEN, SOUTH CAROLINA

(TEL & TEL ADDRESS, AUGUSTA, GA.)

XPLOSIVES DEPARTMENT

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CC:		J. Wende-J. W. Babcock	Croach, Wilm.
	-	Thayer	DPST65-311
		Bellas	
		Krues1, SRP	DE84 005571
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		Jacobsen-W. R.	Kritz
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		Morris	
		Ice-E. L. Albe	
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REA-M16-8

June 7, 1965

J. D. ELLETT, MANAGER SAVANNAH RIVER PLANT

TTENTION: L. W. FOX

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EVALUATION OF MACRORETICULAR ANION EXCHANGE RESIN - RTA-893-R



Introduction

Macroreticular anion exchange resin, which is characterized by a rigid bore structure, was evaluated for use in the reactor purification beionizers in response to RTA-893-R. Macroreticular resins are reported to possess greater physical stability and resistance to oxidation than hormal gel-type anion exchange resins, but they are also slightly more rostly and have a slightly lower volume exchange capacity. Because the tervice life of 100-Area deionizers is limited by radiolytic degradation of the resins¹⁰, the primary incentive for using the macroreticular resin would have to come from an improvement in radiation resistance over the resins currently in use.

In this work the stability of macroreticular resin Amberlite[®] IRA-900 s gamma radiation was determined and compared with that of Amberlite[®] FA-400². An experimental resin volunteered by Ionac Chemical Company is possibly radiation resistant was also examined.

Technical Progress Report, Atomic Energy Division, September 1963. DP-63-1-9, pp. I-20 ff. (Secret)

E. W. Baumann. "Effects of Gamma Radiation on Individual and Mixed Ion Exchange Resins". DPST-64-564. December 31, 1964. Trademark of Rohm and Haas Company, Philadelphia, Pa. DESTRIBUTION OF THIS DOCUMENT IS UMLIMITED

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Summary

The macroreticular anion exchange resin, Amberlite[®] IRA-900-OH, and an experimental resin from Ionac Chemical Company were irradiated by a ¹⁰Co source to doses of 5×10^7 rad and 10^8 rad. These doses approximate or exceed the dose encountered by the deionizer resins in 100-Area service.

The loss in exchange capacity and the volume changes of Amberlite[®] IRA-900 on irradiation were similar to those found previously for Amberlite[®] IRA-400. The Ionac experimental resin, which had a considerably lower initial exchange capacity, likewise was not stable to radiation.

It is concluded that Amberlite[®] IRA-900 offers no advantage over Amberlite[®] IRA-400 for 100-Area purification service. Since there is little justifiestion for further evaluation of macroreticular resin for 100-Area use, the present work completes RTA-893-R.

Mecussion

Background

The reactor deionizer resins accumulate a radiation dose during service that is sufficient to cause a loss of exchange capacity which reduces the service life of the deionizer bed. The anion resin component of the mixed bed limits deionizer lifetime not only because it is preferentially depleted by the acidic moderator but also because it has a lower resistance to radiolytic degradation than the cation resin. Therefore an anion exchange resin with better radiation stability would be of value for use in the 100-Area leionizers.

The recently marketed macroreticular ion exchange resins seemed to offer possible increased resin stability as well as other advantages. In contrast to the conventional gel resins, which shrink and swell, macroreticular resins have an open structure with well-defined pores. According to the anufacturer they also are less susceptible to attrition and oxidation than el resins and are more resistant to organic fouling. The pore structure hould provide greater absorption of particulate matter and improve the apability of the purification system to remove long-lived activities.

Resins Tested

he resins tested for radiation stability were Amberlite[®] IBA-900 (OH Mana), Ionac XAX-1071 (OH form) and a mixture of Amberlite[®] 200 (H form) and the Amberlite[®] IRA-900.

berlite[®] IRA-900 is the macroreticular counterpart of the gel resin berlite[®] IRA-400, the anion exchange resin currently used in 100-Area rification systems. Both resins have quaternary ammonium exchange groups the trimethylamine type.

• experimental resins were supplied by Ionac Chemical Company, as saibly radiation resistant. One of these, Ionac XAX-1072, apparently at exchange sites as it was converted to the hydroxyl form and it was considered further. The other resin, Ionac XAX-1071, was carried ough the irradiation testing procedure. The nature of this experital resin was not established; the infrared absorption spectrum was irely different from the macroreticular and gel resins. The exchange paps are probably of the amine type. Fox

1so tested was a 1:1 equivalent mixture of Amberlite[®] IRA-900 and mberlite[®] 200. Amberlite[®] 200 is the macroreticular counterpart of mberlite[®] IR-120, the cation exchange resin currently used. Amberlite[®] of was tested previously and found to have stability similar to the gel sin; neither resin undergoes appreciable damage in the dose range of interest.

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Experimental Procedure

he anion exchange resins were converted to the hydroxyl form with 10% NON solution and rinsed free of excess caustic. Exchange capacities are determined by placing the resin in a 20% NaCl solution and titrating the alkalinity released with 0.1 N HCl. The resins, now in the Cl form, are dried and weighed.

aberlite[®] IRA-900, Ionac XAX-1071, and a 1:1 equivalent mixture of a berlite[®] 200 and IRA-900 were irradiated in the water-saturated state it a ⁶⁰Co source.

fter a dose of approximately 5×10^7 rad, the resins were rinsed free rester-soluble products and a portion of each anion resin was taken or exchange capacity determination. The remainder of the resin was radiated a second time, for a total dose of approximately 10^8 rad. ain water-soluble products were rinsed out and the exchange capacity r the anion resins determined.

wet packed volume of each resin was determined before and after madiation.

Results

he initial exchange capacities of the resins investigated are presented in Table I for comparison with that of Amberlite[®] IRA-400. The exchange apacity per unit weight is greater for the macroreticular resin than for be gel resin, but the capacity per unit volume is slightly less. The apacity of the Ionac resins was relatively low.

In Figure 1 the loss of initial exchange capacity with gamma dose is seen to be similar for Amberlite[®] IRA-900 and Amberlite[®] IRA-400². The Ionac tesin loses a greater fraction of its capacity than do the Amberlite resins.

lso in Figure 1 are shown changes in volume of the resin on irradiation. Solume changes for macroreticular resins and their mixtures are similar those of the gel resins.

ble II gives information about the water-soluble material from the gradiated resins. Some of the exchange capacity loss of the anion sins was accounted for as water-soluble alkalinity, which could be itrated. As with gel resins, little soluble material was released from the mixed resin. nature of the leach water from the macroreticular anion resin was ifferent from that for gel resins, where the water was brown and murky if it contained suspended polymer. The turbidity in the macroreticular each was mostly filterable and was due to resin fines.

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F. W. Baumann

Elizabeth W. Baumann Analytical Chemistry Division

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L. W. Fox

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TABLE I

Initial Exchange Capacity of Resins

	Exchange Capa	change Capacity	
	meq/g anhydrous resin	meq/ml	Type of Resin
pberlite IRA-400-Cl	(3.3)*	(1.2)	gel
eberlite IRA-900-Cl	3.8 (4.4)	0.9 (1.0)	macroreticular
onac XAX-1071-C1	1.8		experimental; not knowwn
onac XAX-1072-Cl	0.5	* = *	experimental; not known

values in parentheses are from trade literature; others are experimental.

TABLE II

Characteristics of Water-Soluble Material from Resins

Properties of Supernate pH 6; Faintly tan, turbid,	Water-Soluble <u>Alkalinity</u>
pH 6: Faintly tan. turbid.	N.
pH 6: Faintly tan. turbid.	•
foamy; no odor	
pH 11; colorless, turbid, foamy; strong NH ₃ odor	0.7
pH ll; colorless, very clear; faint NH ₃ gdor	0.6
rad total):	
pH 6.5; slightly yellow, turbid foamy, faint NH ₈ odor	l,
pH 11; colorless, turbid, fcamy; strong NH, adam	0.3
pH 9; colorless, wery clear;	0.7
no odor	0.7
	pH 11; colorless, turbid, foamy; strong NH ₃ odor pH 11; colorless, very clear; faint NH ₃ odor rad total): pH 6.5; slightly yellow, turbid foamy, faint NH ₃ odor pH 11; colorless, turbid, foamy; strong NH ₄ odor pH 9; colorless, wery clear; pH 9; colorless, very clear;

