

September 23, 1993

Mr. Matt Nowland
C.R. Bard, Inc.
200 Ames Pond Drive
Tewksbury, MA 01876

SUBJECT:	<u>MAF Number Assigned:</u>	<u>MAF-489</u>
	<u>Date of Submission:</u>	<u>May 27, 1992</u>
	<u>Title of Submission</u>	<u>Geon Resin 30</u>
	<u>Holder of Submission:</u>	<u>BFGoodrich</u>
	<u>Submitted By:</u>	<u>BFGoodrich</u>
	<u>Agent(s):</u>	<u>None</u>

Dear Mr. Nowland:

This letter is the FDA's authority to refer to MAF 489 for the information contained therein on Geon Resin 30 in the review of any of your premarket approval applications (PMA's), investigational device exemptions (IDE's), premarket notification submissions, reclassification petitions, color additive petitions, or other submissions to the FDA. You should include the original copy of this letter in the original copy of your submission to the FDA in order that your submission will not be found deficient for the lack of authorization and to expedite processing.

Any Geon Resin 30 supplied to you for this purpose will comply with the product specifications contained in MAF 489.

Sincerely,

THE BFGOODRICH COMPANY

Connie N. Dillon
Regulatory and Data
System Specialist

bcc: Tom W. Williams (OTM-d/5-375)

BFG16258

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