

/n

OFFICE OF NAVAL RESEARCH Contract: N00014-85-K-0222 Work Unit: 4327-555 Scientific Officer: Dr. Richard S. Miller

Technical Report No. 28

MATRIX CRACKING INITIATED BY FIBER BREAKS IN MODEL COMPOSITES

by

A. N. Gent and C. Wang

Institute of Polymer Engineering The University of Akron Akron, Ohio, 44325-0301

December 1991

Reproduction in whole or in part is permitted for any purpose of the United States Government Approved for public release; distribution unrestricted

1930 000

9]



SECURITY CLASSIFICATION OF THIS PAGE (When Date Entered)					
REPORT DOCUMENTATION PAGE	READ INSTRUCTIONS BEFORE COMPLETING FORM				
1. REPORT NUMBER 2. GOVT ACCESSION NO.	3. RECIPIENT'S CATALOG NUMBER				
Technical Report No. 28					
4. TITLE (and Sublille)	5. TYPE OF REPORT & PERIOD COVERED				
MATRIX CRACKING INITIATED BY FIBER BREAKS	Technical Report				
IN MODEL COMPOSITES	6. PERFORMING ORG. REPORT NUMBER				
7. AUTHOR(+)	8. CONTRACT CR GRANT NUMBER(*)				
A. N. Gent and C. Wang	N00014-85-K-0222				
9. PERFORMING ORGANIZATION NAME AND ADDRESS	19. PROGRAM ELEMENT, PROJECT, TASK AREA & WORK UNIT NUMBERS				
The University of Akron Akron, OH 44325-0301.	4327-555				
11. CONTROLLING OFFICE NAME AND ADDRESS	12. REPORT DATE				
Office of Naval Research	December 1991				
Power Program Arlington VA 22217-5000	13. HUMBER OF PAGES				
14. MONITORING AGENCY NAME & ADDRESS(If different from Controlling Office)	15. SECURITY CLASS. (of this report)				
	UNCLASSIFIED				
	154. DECLASSIFICATION/DOWNGRADING SCHEDULE				
Approved for public release; distribution unrestricted.					
17. DISTRIBUTION STATEMENT (of the abetract entered in Block 20, 11 different from Report)					
18. SUPPLEMENTARY NOTES					
Submitted to: Journal of Materials Science					
19. KEY WORDS (Continue on reverse side if necessary and identify by block number)					
Cohesive Fracture, Composites, Crack Growth, Energy Release Rate, Fiber Reinforcement, Fracture Mechanics, Resin Cracking, Strength					
20. ATSTRACT (Continue on reverse elde if necessary and identity by block manber)					
Fracture of resin in a composite material can be initiated by a tensile break in a fiber. This process has been investigated for a simple model composite, consisting of two inextensible rods placed along the axis of a cylindrical elastic block and touching in the center. The rods represent a broken fiber. Energy release rates G were calculated by FEM for a circular crack growing					
outwards from the point where the rod ends separated as they were					

Ъ.

.

SECURITY CLASSIFICATION OF THIS PAGE ("Then Data Entered)

pulled apart. Results are compared with experimental observations on cracking of a silicone rubber cylinder containing two steel It was found that a crack grew outwards under increasing rods. load until its radius reached a certain size, approximately half way to the surface of the resin cylinder. At this point G reached a minimum value and then increased. Simultaneously, the crack accelerated and the sample broke. Forces required to propagate the crack were successfully predicted by linear elastic fracture mechanics at all stages of crack growth and for a wide range of fiber and sample radii. In particular, good agreement was obtained with the maximum force that the model system could support, i.e., the breaking load. When the sample was surrounded by a rigid tube, representing neighboring fibers surrounding the broken one, growth of a crack required an increasing load at all stages. The sample finally fractured when the broken fiber pulled out with resin still attached to it. Application of these results to unidirectional fiber-reinforced materials is discussed.

S/N 0102- LF- 014- 6601

SECURITY CLASSIFICATION OF THIS PAGE ("Then Date Entered)

Matrix Cracking Initiated by Fiber Breaks in Model Composites

A. N. Gent and C. Wang

Institute of Polymer Engineering

The University of Akron, Akron, OH 44325-0301, U.S.A.

Abstract

Fracture of resin in a composite material can be initiated by a tensile break in a fiber. This process has been investigated for a simple model composite, consisting of two inextensible rods placed along the axis of a cylindrical elastic block and touching in the center. The rods represent a broken fiber. Energy release rates G were calculated by FEM for a circular crack growing outwards from the point where the rod ends separated as they were pulled apart. Results are compared with experimental observations on cracking of a silicone rubber cylinder containing two steel rods. It was found that a crack grew outwards under increasing load until its radius reached a certain size, approximately half way to the surface of the resin cylinder. At this point G reached a minimum value and then increased. Simultaneously, the crack accelerated and the sample broke. Forces required to propagate the crack were successfully predicted by linear elastic fracture mechanics at all stages of crack growth and for a wide range of fiber and sample radii. In particular, good agreement was obtained with the maximum force

that the model system could support, i.e., the breaking load. When the sample was surrounded by a rigid tube, representing neighboring fibers surrounding the broken one, growth of a crack required an increasing load at all stages. The sample finally fractured when the broken fiber pulled out with resin still attached to it. Application of these results to unidirectional fiber-reinforced materials is discussed.



Accesion For				
NTIS DTIC Udouu Justific	CRA&I TAB ounced ation			
By Dist ibtion/				
Availability Corles				
Dist	Avail and for Special			
A-1				

1. Introduction

Fiber-reinforced composites are light-weight and strong materials, with several possible modes of rupture. At present there is no satisfactory theory relating strength of composites to properties of the constituent materials, even in the simplest cases. Two models of a composite with continuous fiber reinforcement are employed here to study the consequences of a fiber break. The first model is shown in Figure 1a. Two identical steel rods were embedded along the axis of a cylindrical block of silicone rubber resin, with their ends in They were treated with a primer to give good bonding contact. to the silicone rubber. Because of the great difference in Young's modulus of steel and silicone resin, the rods can be considered to be inextensible in comparison with the elastic matrix. Thus, when the sample is put under a tensile load all of the strain energy is stored in the matrix resin, assumed for simplicity to be linearly-elastic.

The second model was used to investigate the effect of neighboring fibers on a crack propagating outwards from the initial break. It is shown in Figure 1b. It differs from the first model by the addition of a rigid tube enclosing the resin cylinder and bonded to it. The sample thus consists of a tube of silicone resin, bonded to a broken rigid fiber at its center, and to a concentric rigid tube surrounding it. The external

tube represents the ring of parallel fibers surrounding a given fiber in a multi-fiber composite. Strong bonding between the resin, the central fiber, and the external tube, ensured that adhesive failures did not occur in the experiments.

A Griffith fracture criterion is applied, as follows (1):

$$- \partial W/\partial A|_{\rho} \geq G_{C}$$
(1)

where \underline{W} is elastic strain energy stored in the system (in the present case, in the resin only) \underline{A} is crack area and \underline{G}_{C} is fracture energy of the resin. The derivative is evaluated at constant length $\underline{\ell}$, and hence displacement, so that the applied force does no further work as the crack grows. Thus, energy required for fracture is obtained solely from elastically-stored energy. The left-hand side of Equation 1, termed the strain energy release rate, is denoted \underline{G} hereafter.

Equation 1 gives a necessary condition for a crack to propagate: the rate of release of strain energy must reach a critical value. In some cases crack growth will be stable, i.e., the applied load must be increased to grow the crack further. In other cases growth is unstable, resulting in catastrophic failure. The criterion for stable growth is:

$$dG/dA \leq 0$$
, at $G = G_c$. (2)

When a fiber breaks, two types of failure may ensue. A debond may propagate along the fiber/matrix interface in the form of a cylindrical crack running in the fiber direction, or a planar (circular) crack may grow outwards into the matrix material at right angles to the fiber. Which mode of failure takes place depends on the geometry and mode of loading of the specimen and the relative strength of adhesion compared to strength of the resin. If adhesion is relatively weak, a crack will propagate along the interface as a debond. On the other hand, if adhesion is strong, then a perpendicular crack will grow outwards into the resin - which is the case considered here.

Debonding of fibers, "pull-out", and repeated fracture of fibers as a result of progressive failure of the bond between fiber and resin, have been examined elsewhere (2-5). A critical force is required to initiate an interfacial crack at the fiber end. Values were calculated from a Griffith fracture criterion, Equation 1, with \underline{G}_{c} replaced by a fracture energy for rupture of the interfacial bond , denoted \underline{G}_{a} (6). (Values of \underline{G}_{a} are generally smaller than \underline{G}_{c} .)

As a fiber is debonded, frictional resistance to pull-out increases and the force needed to propagate the debond increases continuously with the length debonded (3). This effect can lead to repeated breaking of the fiber, termed "fragmentation". The frictional effect has been minimized in other work by using "push-out", instead of "pull-out", mechanics (4). In practice,

the main parameter which determines whether debonding will occur or not in a given composite is \underline{G}_a , the energy required to debond unit area of fiber-matrix interface.

Compared with fiber pull-out, the mechanics of resin cracking have received little attention. Results for two extreme cases are well-known. Growth of a small penny-shaped crack of radius <u>a</u> in a homogeneous material has been analyzed by Sack (7). The critical far-field tension stress is:

$$\sigma^2 = \pi E G_{c} / 3a. \tag{3}$$

The corresponding problem for a circular debond located at the interface between an elastic half-space and a rigid substrate was solved by Mossakovskii and Rybka(8). The result is similar:

$$\sigma^2 = 2\pi E G_{\rm C} / 3a, \qquad (4)$$

the additional factor of 2 arising from the presence of elastically-stored energy in only one-half of the composite.

These examples can be regarded as extreme cases of matrix cracking (2). They both predict catastrophic growth of the initial crack when the critical condition is achieved, because the stress required to propagate a crack decreases as the crack radius increases. The more complex case when a broken fiber is the initiator of a crack has been considered by Mullin et al.(9).

They considered a single inextensible fiber embedded in an elasto-plastic resin and found that either one circular crack, two inclined-conical cracks, both, or neither, formed at the break and grew into the matrix. Which process occurred depended on the particular combination of fiber and matrix considered. Mahishi <u>et al</u> used a finite element model to examine crack propagation in an elasto-plastic matrix material (10,11). They found that the stress intensity factor increased with crack radius up to a certain point. Thereafter, unstable crack growth was predicted to occur.

A greater release of strain energy may be obtained for a cylindrical crack that propagates along the fiber, in the resin but as close to the fiber as possible, rather than for a circular crack growing at right angles to the fiber. This is probably the case when the fiber radius is small in comparison to the radius of the cylinder of resin (2). We are concerned here only with growth of circular cracks, as observed when the fiber radius is relatively large compared to that of the resin cylinder surrounding it and the bonding between fiber and resin is strong.

Experimental measurements of crack growth and breaking force have been carried out for the simple two-rod model, using various ratios of rod radius to resin cylinder radius. The results are compared with theoretical predictions from FEM calculations. The objectives of the study are to elucidate the principal factors affecting resin cracking initiated by a broken fiber and

to predict the strength of a model fiber composite when this is the mode of failure. The analysis is also relevant to the widely-used "fragmentation test" for adhesion between resin and fiber (12-15).

2. Stress intensity factors and strain energy release rate

The stress field around a broken fiber end is quite complex and analytical solutions are rather difficult to obtain. Finite element numerical techniques have been widely used, therefore, to determine crack tip stress fields and stress intensity factors. In some cases special crack tip elements were employed to deal with the stress singularity at the crack tip (16). But stress intensity factors can also be derived from energy release rates, as described below. Finite element methods were used here to derive values of energy release rate \underline{G} for various sizes of crack in the resin, from the effect of a crack on the sample compliance. Dixon (17) showed that the stress field in the immediate vicinity of the crack tip makes relatively little contribution to the overall displacement of the system, and hence to the compliance, so that excessive refinement of the mesh at the crack tip is not necessary. This is the principal advantage of energy methods as employed here.

The first step is to calculate the elastic compliance \underline{C} of specimens containing cracks of increasing radius \underline{a} . Strain energy release rates associated with the creation of new crack

surface are then obtained by numerical differentiation of the relation between compliance and crack area <u>A</u> (18):

$$G = (F^2/2)\partial C/\partial A = (F^2/4\pi a)\partial C/\partial a.$$
 (5)

The following procedure was employed:

First, for a given value of crack radius <u>a</u>, the displacement δ was calculated for a given tensile force <u>F</u> applied to the outer ends of the rods. In this way the compliance <u>C</u> was determined, $\underline{C} = \delta/\underline{F}$. Then, by incrementing the crack radius, the rate of change of compliance with crack radius was evaluated and hence <u>G</u> from Equation 5. Finally values of the applied force <u>F</u> at which a given crack would grow were calculated from Equation 5 by replacing the value of <u>G</u> with its critical value, the fracture energy <u>G</u> of the resin. Hence:

$$F^{2} = 4\pi aG_{c}/(\partial C/\partial a).$$
 (6)

In linear elastic fracture mechanics, the relation between energy release rate and stress intensity factor <u>K</u> in plane strain is (18)

$$K^{2} = EG/(1 - v^{2})$$
(7)

where \underline{E} is Young's modulus and ν is Poisson's ratio of the resin.

The dimensionless stress intensity factor \underline{Y} is defined by (18)

$$K = Y \sigma \sqrt{\pi a}$$
(8)

where σ is the far-field stress, $\sigma = \underline{F}/\pi \underline{R}_0^2$, where \underline{R}_0 is the radius of the resin block. Values of \underline{K} and \underline{Y} can thus be calculated from \underline{G} by means of Equations 7 and 8, and values of breaking force \underline{F}_b obtained on substituting a critical value for \underline{K} , denoted \underline{K}_c , in Equation 8. Thus, exactly equivalent calculations can be carried out in terms of stress intensity \underline{K} and its critical value \underline{K}_c , in place of strain energy release rate \underline{G} , with a critical value \underline{G}_c .

3 Details of experimental procedure and FEM calculations

3.1 Preparation of test-pieces and properties of the resin

Transparent silicone resin (Sylgard S-184, Dow Corning Corporation) containing 10% by weight of curing agent(Sylgard C-184, Dow Corning Corporation) was used as matrix material, and flat-ended steel rods as fibers. To obtain good bonding to the resin the cylindrical surfaces of the rods were first polished with fine emery paper, cleaned with isopropanol and dried. They were then immersed in Primer 92-023 (Dow Corning Corporation) for 24 hours. A small amount of partially-cured resin was used to glue two rods together end-to-end. The joined rods were then carefully placed along the axis of a plastic tube,

used as a mold for casting the cylinder of resin. Uncured silicone resin was poured into the tube to surround and encapsulate the steel rods. Curing was effected by heating the assembly for 12 hours at 110° C. Samples were made with different combinations of rod radius \underline{R}_i and external radius \underline{R}_o of the resin cylinder: $\underline{R}_i = 0.435$ to 1.16 mm; $\underline{R}_o = 3.0$ and 5.6 mm. All samples had a length \underline{L} of 50 mm.

For the second set of experiments, glass tubes were bonded to the external surface of the cured resin cylinder to simulate the constraints due to neighboring fibers. The internal and external radii of the glass tubes were 3 and 4 mm, respectively. Strong bonding between the silicone resin and glass was obtained by immersing the tubes in a 50 % solution of adhesive primer in heptane for 12 hours and then drying them at room temperature before use. Rods, glued together, were placed along the axis of the tube, which was then filled with uncured resin. Curing was effected at room temperature for 7 days to avoid cracking caused by shrinkage stresses, either at the fiber/resin or tube/resin interface, that was encountered on cooling from an elevated curing temperature. Typical dimensions of a specimen were: rod radius $\underline{R_i} = 0.435$ mm, external radius $\underline{R_o}$ of the resin cylinder = 3.0 mm, overall length \underline{L} of cylinder = 50 mm.

Tensile stress-strain relations for the cured silicone resins were found to be quite linear up to about 30 percent strain. Young's modulus \underline{E} was determined from the slope at

small strains to be 2.8 MPa for samples cured at 110° C and 0.87 MPa for samples cured at room temperature.

Fracture energies \underline{G}_{C} of silicone resin were determined at different rates of tear propagation, from measurements of the average force required to propagate a tear. The results are plotted in Figure 2 against the rate of tearing. Results for both curing conditions were approximately the same, even though the values of tensile modulus were quite different. As for other viscoelastic materials there is a considerable effect of tear rate, the value of \underline{G}_{C} increasing from about 180 J/m² at a rate of 0.8 µm/s to about 330 J/m² at a rate of 2 mm/s. In experiments with both models of a composite, crack growth took place at rates lying in the relatively narrow range: 5 to 25 µm/s. Thus, the relevant value of \underline{G}_{C} in the fracture experiments was 200 to 230 J/m².

3.2 Measurement of crack size and sample compliance

Specimens were loaded in tension using an Instron tensile testing machine equipped with a microscope, video camera and recorder to study crack growth as the applied tensile load was increased continuously. When the steel rods were pulled apart, a crack initiated at the point where they were initially in contact, and grew transversely across the matrix. Its radius <u>a</u> was measured as the distance from the axis of the rods to the crack tip. The relation between load and displacement was

recorded simultaneously, and the compliance of a specimen containing a crack of a given radius was derived from the chord of the displacement-load relation at the appropriate displacement, as shown in Figure 3. Compliances for various crack radii were obtained in this way in a single experiment. A relatively slow rate of stretching, of 8 μ m/s, was employed. All measurements were carried out at room temperature, about 23°c.

Due to its cylindrical surface, the transparent silicone resin served as a lens and magnified the size of the crack. The degree of magnification was approximately constant as the crack grew, given by the refractive index of the resin, 1.41X (see Appendix 1). Corrections were made to the observed crack radius to allow for this magnification.

In contrast, the applied tensile stress, enhanced somewhat by stress concentrations near the crack tip, caused the resin cylinder to undergo Poissonian contraction and the crack became smaller in radius than in the unstressed state. A correction for the amount of contraction was deduced by FEM, as described in Appendix II. Generally speaking, the effect of magnification on the apparent crack size was about + 40% and that of Poissonian contraction was about - 10%.

Figure 3 shows a typical force-displacement relation for a model composite specimen without external constraints, loaded in tension. It is linear initially, up to the point B.

Non-linearity starting at B is attributed to crack propagation. Thus, the load at B, about 3.0 N, is taken as the force required to initiate growth of a crack with a radius given by the rod radius. Then, as the crack grew, the compliance of the specimen increased continuously.

Linear elastic behavior of a cracked specimen was verified by stretching a sample with a large-diameter crack already developed at the rod end, and then unloading the specimen before the crack started to grow further. A linear relation was obtained between load and displacement for this cracked specimen, with virtually no hysteresis between loading and unloading relations.

3.3 <u>Pre-stress due to thermal contraction</u>

A curious feature of the experimentally-determined relations between load and displacement was a small offset in the initial portion of the force-displacement curve (region O-A in Figure 3). In effect, a certain force was required to pull the rods apart initially. This is attributed to a prestress set up by differential thermal contraction between the rods and resin as the specimen was cooled down from curing temperature to room temperature. As a result, a shrinkage stress developed in the resin, pulling the rods together. The shrinkage force could be estimated by extrapolating the linear behavior observed at small displacements (region A-B) back to the load axis. A residual force of about 0.9 N was deduced in this way, approximately 8% of

the maximum load required to break the sample. Figure 4 shows the variation of residual force with test temperature. It decreased continuously, becoming zero at 100°C, close to the curing temperature, confirming that the source of this anomalous feature of the load-displacement relation at room temperature was a compressive pre-load caused by thermal shrinkage of the resin. In contrast, there was no thermal contraction in the second composite model because the resin was cured at room temperature in that case.

3.4 Numerical calculations

An ADINA FEM program (20) was used to calculate compliances of a specimen with different crack radii. Eight-noded quadrilateral axisymmetric elements with four integration points were used. The mesh is shown in Figure 5. Because of the symmetry of the system, only one-half of the specimen was modeled. Fifteen elements were employed in the radial direction and thirty elements in the axial direction, between the rod end and upper end of the cylinder. For the second model, with external constraints imposed by the outer rigid tube, all nodes on the outer surface, radius \underline{R}_0 , were fixed. The maximum aspect ratio of mesh elements was about 4.5.

Both rods and resin were assumed to be linearly-elastic and nearly incompressible, with Poisson's ratio of 0.4999. To make the rods effectively rigid, Young's modulus was made 10⁹ times

that of resin. Perfect bonding was assumed to exist between the rods and resin (and between the resin and external tube, in the second model). Different radii \underline{R}_i and \underline{R}_o were employed, with the ratio $\underline{R}_i/\underline{R}_o$ ranging from 0.053 to 0.8. The ratio of cylinder length \underline{L} to resin radius \underline{R}_o was chosen to be long enough to minimize end effects and still give adequate accuracy for determining small changes in compliance with crack radius. Preliminary results indicated that a value of 16.6 was satisfactory in this respect; it was used in all the cases reported here. Most of the calculations were carried out for a representative specimen with dimensions: $\underline{R}_i = 0.435 \text{ mm}$, $\underline{R}_o = 3 \text{ mm}$, $\underline{L} = 50 \text{ mm}$.

3.5 Test of validity of FEM results

The validity of the FEM model was examined by solving a simple case for which an analytical solution is already known. A penny-shaped crack in the center of a homogeneous elastic cylinder under tension has been widely investigated (21). An approximate solution for a dimensionless stress intensity factor, denoted Y' and defined by the relation:

$$K = Y' a^{1/2} [1 - (a/R)]^{1/2} [F/\pi (R^2 - a^2)]$$
(9)

was obtained by cubic interpolation (21). It takes the form:

$$Y' = (2/\pi) [1 + (1/2) (a/R) - (5/8) (a/R)^{2}] + 0.268 (a/R)^{3}.$$
 (10)

When the crack size is much smaller than the radius <u>R</u> of the cylinder, the value of <u>Y</u>' (now equal to <u>Y</u>, Equation 8) is $2/\pi$, in agreement with Sack's result, Equation 3.

Values of the compliance <u>C</u> and its derivative $\partial C/\partial A$ with crack area were calculated using the present finite element model. Values of <u>Y</u>' calculated from them by means of Equations 5, 7 and 9 are compared with values from Equation 10 in Figure 6. Agreement is quite good, within 5 percent over most of the range of crack radius. Discrepancies for extremely small and extremely large cracks are attributed to the use of only 15 elements to represent the entire radial distance for crack growth. Thus, small cracks were represented by only one or two elements and, for large cracks, only one or two elements remained unbroken. FEM results would not be expected to be highly accurate under these circumstances.

4. Results and discussion

4.1 <u>Results for an unrestrained sample (model 1)</u>

4.1.1 Compliance as a function of crack size

Values of compliance were calculated as a function of crack radius for various values of rod radius \underline{R}_i and sample radius \underline{R}_o . Results for specimens with a external radius \underline{R}_o of 3 mm are

plotted in Figure 7. The compliance increased with increasing crack size, in an approximately linear way, and then more rapidly as the dimensionless crack radius $(\underline{a}-\underline{R}_{1})/(\underline{R}_{0}-\underline{R}_{1})$ exceeded about 0.6. And the larger the radius of the enclosed rods, the lower was the compliance. Similar relations were obtained for samples with different external radius \underline{R}_{0} . As expected, the compliance was smaller for samples of larger radius.

In the experiments, a circular crack was found to grow symmetrically, under increasing load, until it reached a certain size, about half way to the edge of the sample. The load then decreased slightly and catastrophic failure followed immediately. Stages in the growth of a typical crack are shown in Figure 8. Values of sample compliance were recorded continuously as the crack grew, until the sample fractured. Figure 9 shows a comparison of compliance results from FEM and from direct measurement. Good agreement is seen to hold where comparison is possible, i.e., until catastrophic failure intervened.

4.1.2 Energy release rates and breaking forces:

Rates of change of compliance with crack area $\partial \underline{C}/\partial \underline{A}$ were calculated by a central difference method using three successive values of \underline{C} as a function of crack radius. From these results, values of \underline{G} for a unit applied tensile load were obtained using Equation 5. They are plotted in Figure 10 as a function of dimensionless crack radius, for samples with an external radius

of 3 mm and different rod radii. Initially, <u>G</u> decreases with increasing crack size and then passes through a broad shallow minimum at a dimensionless crack radius of about 0.5. Similar results were obtained for different rod radii and with other sample radii.

Corresponding values of force <u>F</u> required to propagate a crack were then calculated from Equation 6, using the appropriate value of fracture energy \underline{G}_{c} of 230 J/m_{J}^{2} for the silicone resin used in the experiments. They are represented in Figure 11 by a continuous curve. As a crack starts to grow from its initial size, given by the rod radius, an increasing load is seen to be needed to propagate it. Thus, crack growth is initially stable. But after the required force passes through a maximum (and the energy release rate passes through a minimum), the system becomes unstable and catastrophic failure ensues.

Experimental measurements of crack radius are plotted in Figures 11 and 12 against the applied load, for comparison with calculated values. Good agreement was obtained, both in the general form of the relation for crack propagation force as a function of crack radius, and in the breaking force when the crack reaches a critical size. A similar degree of agreement was obtained with rods and samples of various radii. Thus, the theoretical treatment appears to describe the process of crack growth with considerable success. Experiments could not be continued beyond a dimensionless crack radius of about 0.5

because the samples broke at this point, as predicted.

4.1.3 Stress to initiate crack growth:

The applied force \underline{F}_i and corresponding stress σ_i required to initiate crack growth were estimated using Equation 6. Initial values of \underline{G} were obtained by fitting calculated values to a five-term polynomial function of crack radius and then extrapolating back to a crack radius equal to that of the rod. Values of \underline{F}_i and σ_i obtained in this way are given in Table 1. They <u>increase</u> as the radius of the rods, and hence the radius of the starter crack, increases. This is in striking contrast to an isolated circular crack, Equation 3, where larger cracks require smaller stresses to propagate them. Apparently the presence of rigid bonded rods has a protective effect on cracks formed at their ends.

The effect of sample radius is shown in Table 1 also. For a given rod radius, the stress σ_i required to initiate a crack was found to decrease as the sample radius increased, although the force, of course, increased.

4.1.4 Breaking stress:

Values of breaking force \underline{F}_{b} and breaking stress σ_{b} were calculated from minimum values of <u>G</u>, using Equation 6. The results are given in Table 1. When the rod radius was smaller than about 0.5 \underline{R}_{o} , the minimum value of <u>G</u> was found to be

approximately constant, and was reached at a critical crack radius \underline{a}_{C} of about $(\underline{R}_{i} + \underline{R}_{o})/2$, i. e., when the crack was about half-way to the edge of the sample. On the other hand, when the rod radius was larger than about 0.5 \underline{R}_{o} , the minimum value of \underline{G} was larger and occurred at a smaller crack radius. In other words, when the rod radius is smaller than about one-half of the sample radius, fracture would be expected at a dimensionless crack radius of about 0.5 at a constant force, independent of rod radius, but when the rod radius is larger than this, fracture is predicted to occur at an earlier stage of crack growth and at a smaller force. In contrast, the force required to start a crack growing depends directly upon its initial radius \underline{R}_i , as shown in Table 1: the larger the value of \underline{R}_i , the larger the force needed to initiate crack growth. Experimental results were in good agreement with these predictions in all cases.

Samples with a larger external radius were found to break at a lower stress, even though the breaking force was higher, Table 1. In fact, the breaking stress was somewhat smaller (0.55X to 0.7X) than the corresponding value for an isolated circular crack of the same radius in a homogeneous solid, Table 1. Thus, although embedded rods strengthen a sample initially and prevent small cracks from growing catastrophically, they have a slight weakening effect when the crack reaches its critical size.

It should be noted that the stress to fracture a composite depends on both \underline{R}_i and \underline{R}_o separately, and not on the ratio $\underline{R}_i/\underline{R}_o$.

For example, if we compare a sample with $\underline{R}_i = 2.4$ mm and $\underline{R}_o = 3.0$ mm with another sample with $\underline{R}_i = 0.435$ mm and $\underline{R}_o = 0.545$ mm, both composites have the same $\underline{R}_i / \underline{R}_o$ ratio but the breaking stresses are different: 0.29 and 0.62 N/mm².

All of the FEM results indicate the rather surprising feature that a crack will grow from a fiber break in a stable way, requiring an increasing tensile load, until it is about half way to the edge of the specimen. This is in marked contrast to a crack in a homogeneous medium or at the horizontal interface between an elastic and a rigid material, when all cracks are, in principle, unstable.

In a fiber fragmentation test, cracking of the resin is sometimes observed, originating at the broken fiber ends (13-15). Diameters of these resin cracks can be several hundred times larger than the fiber diameter without fracturing the specimen (13). This feature is consistent with the observations of crack growth and crack stability reported here.

4.2 Effect of neighboring fibers (model 2)

A rigid cylindrical tube bonded to the external surface of the matrix is used here to represent neighboring fibers surrounding a broken fiber. Calculated compliances for different values of rod radius \underline{R}_i are plotted in Figure 13 as a function of dimensionless crack radius. When the dimensionless crack radius is greater than about 0.5, the compliance is seen to

reach a substantially constant value. Corresponding values of the energy release rate <u>G</u>, obtained as before from the rate of change of compliance with crack area, are plotted in Figure 14. They are seen to decrease over the entire range of crack radius. Thus, stable crack growth would be expected for all sizes of crack, until the crack reaches the surface of the externally-restraining tube. Addition of the rigid tube has thus made crack growth more difficult and there is now no tendency for catastrophic growth to occur.

Force-displacement relations were determined experimentally for samples with a rigid tube bonded to the external surface of the resin. Forces due to thermal contraction were absent and, because of the external restraint, Poissonian contraction did not occur. Thus, only the effect of optical magnification needed to be taken into account in determining crack radii. The refractive index of the glass tube was assumed to be the same as that of the silicone resin.

Cracks could be observed up to a radius of 2.7 mm in a sample with resin radius of 3.0 mm, corresponding to a dimensionless crack radius of 0.9, before internal reflection prevented further viewing. Crack growth was stable up to the limit of observation. A comparison is made in Figure 15 between the calculated relation for crack radius as a function of applied force, represented by the full curve, and experimental measurements. Good agreement is seen to obtain over the entire

range of crack radius, using a value for resin fracture energy \underline{G}_{C} of 230 J/m² as indicated by direct measurements on the resin.

Crack growth began at an applied force of 9.5 N, close to the value, 8.4 N, obtained by extrapolating FEM results to the initial crack size. Further crack growth required a larger Lorce. It is clear that the presence of the external tube, representing neighboring fibers around a broken fiber, tends to make resin cracking more difficult and prevent it from becoming catastrophic.

Thus, both the form of the crack growth relation and the actual values of crack driving force appear to be predicted successfully for the second model, as well as for the first.

The fiber content of the composite is represented in model 2 by the ratio of cross-sectional areas of fiber and sample, $\left(\frac{R_i}{R_0}\right)^2$. As this ratio was increased, the force required to propagate a crack increased, Table 2.

5. Conclusions

In general, materials fail when the energy release rate of the system reaches a critical value, \underline{G}_{C} . This criterion is widely used in linear elastic fracture mechanics. If $\underline{dG/dA}$ is negative, an increased stress is needed to maintain crack growth. This results in stable crack growth. It has been shown that a circular crack, created by fiber fracture in a single-fiber model composite, is stable in this way until it reaches a relatively large radius, about half-way to the surface of the cylindrical

resin block containing the fiber. Experimental observations are in agreement with these conclusions. The force required to propagate a crack was found to be in good agreement with values predicted by linear elastic fracture mechanics at all stages of crack growth. In particular, the maximum force that the system could support - the breaking load - was predicted successfully for a range of dimensions. On the other hand, for a model surrounded by a rigid tube representing neighboring fibers in a closely-packed fiber composite, a crack growing outwards from a broken fiber was predicted to be stable at all stages of growth, requiring an increasing force to make it grow further. Again. good agreement was found between calculated and measured values. Eventually, these specimens broke by failure of the resin in shear, after the crack tip had reached the edge of the resin cylinder. A similar analysis of this mode of failure is therefore being attempted.

Acknowledgements

This work is part of a study of the strength of elastomeric composites, supported by the Office of Naval Research (Contract N00014-85-K-0222; Project Officer, Dr. R. S. Miller). Grants-in-aid were kindly provided by Lord Corporation, Westvaco and 3M Company. Assistance with FEA calculations by Professor J. Padovan and Dr. Y.-C. Hwang of these laboratories is gratefully acknowledged also.

References

- A. A. Griffth, Phil. Trans. Roy. Soc. (London) <u>A221</u> (1920)
 163.
- A. N. Gent, G. S. Fielding-Russell, D. I. Livingston and
 D. W. Nicholson, J. Mater. Sci. <u>16</u> (1981) 949.
- 3. A.N. Gent and G.L. Liu, J. Mater. Sci. <u>26</u> (1991) 2467.
- 4. A. N. Gent and S. Y. Kaang, Rubber Chem. Technol. <u>62</u> (1989) 757.
- 5. A. N. Gent and O. H. Yeoh, J. Mater. Sci. <u>17</u> (1982) 1713.
- 6. A. N. Gent, Rubber Chem. Technol. <u>56</u> (1983) 1011.
- 7. R. A. Sack, Proc. Phys. Soc. (Lond.) <u>58</u> (1946) 729.
- 8. V. I. Mossakovskii and M. T. Rybka, Prikl. Mat. Mekh. <u>28</u> (1964) 106.
- 9. J. Mullin, J. M. Berry and A. Gatti, J. Composite Mater.
 <u>2</u> (1968) 82.
- 10. J. M. Mahishi and D. F. Adams, J. Composite Mater. <u>16</u> (1982) 457.
- 11. J. M. Mahishi and D. F. Adams, J. Mater. Sci. 18 (1983) 447.
- 12. L. T. Drzal, M. J. Rich, J. D. Camping and W. J. Park, 35th Annual Technical Conference, 1980, Reinforced Plastics and Composites Institute, (Society of Plastics Industry), <u>20-C</u>, p. 1.
- 13. S. M. Lee and S. Holguin, J. Adhesion <u>31</u> (1990) 91.

- 14. A. T. DiBenedetto and P. J. Lex, Polym. Eng. Sci. <u>29</u> (1989) 543.
- 15. W. D. Bascom and R. M. Jensen, J. Adhesion 19 (1986) 219.
- 16. R. S. Barsoum, Int. J. Numer. Meth. Engng., <u>11</u> (1977) 85.
- 17. J. R. Dixon and L. P. Pook, Nature 224 (1969) **565** 166.
- 18. J. G. Williams, "Fracture Mechanics of Polymers", John Wiley& Sons, New York, 1984, p. 30.
- 19. I. N. Sneddon and M. Lowengrub, "Crack Problems in the Classical Theory of Elasticity", John Wiley & Sons, New York, 1970, p. 138.
- 20. K. J. Bathe, "ADINA: A Finite Element Program for Automatic Dynamic Incremental Non-Linear Analysis", Report No. 82448-1, Massachusetts Institute of Technology, Cambridge, Massachusetts, 1977.
- 21. J. P. Benthem and W. T. Koiter, Chapter 3, "Methods of Analysis and Solutions of Crack Problems", ed. by G. C. Sih, Noordhoff Internatl. Publishing, Leyden, 1973.

Appendix I

Magnification of the apparent crack radius by the cylinder of resin surrounding it can be calculated as follows. As shown in Figure 16, the angles θ_a and θ_b of light rays emerging from the cylinder can be expressed by Snell's law:

$$\sin \theta_{a} = n \left(Y_{o} / R_{o} \right) \tag{11}$$

$$\sin \theta_{b} = n \sin \left[\tan^{-1}(y_{o}/R_{o}) \right]$$
(12)

where <u>n</u> is the refractive index of the resin (n = 1.41), \underline{R}_{o} is the radius of the resin cylinder, and \underline{y}_{o} is the crack radius. Equations for the light path are:

$$y - y_o = - [x - (R_o^2 - y_o^2)]^{1/2} \tan [\theta_a - \sin^{-1}(y_o/R_o)]$$
 (13)

$$y = - (x - R_0) \tan \theta_b \tag{14}$$

The degree of magnification $\underline{y}/\underline{y}_0$ can be determined by inserting the actual crack size \underline{y}_0 into the above equations and solving for \underline{y} . A calibration curve is given in Figure 17, and compared with experimental measurements of the degree of magnification. Over a wide range, the magnification is about 1.4X, approximately the same as the value for small angles, given

by the refractive index of the resin.

Total internal reflection occurred when the incident angle was larger than a critical value. This made it impossible to measure a crack radius larger than about 70% of the cylinder radius.

Appendix II

When a cracked specimen is subjected to a tensile force under which the crack does not propagate but merely opens, the crack is reduced in radius by Poissonian contraction of the entire specimen. The measured crack radius is thus smaller than its actual size in the unstrained state. The higher the stress, the larger the discrepancy. An estimate of the amount of lateral contraction can be obtained by FEM, by determining the relative lateral contractile displacement of the crack tip (δ ') and longitudinal tensile displacement of the upper surface of the cylinder (δ), for unit applied tensile load. A plot of δ ' versus δ is presented in Figure 18 for a sample with rod radius $R_i = 0.435$ mm and resin radius $R_0 = 3.0$ mm. The actual crack radius is then given by

$$a_{actual} = a_{observed} + \delta'$$
. (15)

Table 1. Calculated and measured forces <u>F</u> and stresses <u>g</u> to initiate a crack $(\underline{F}_i, \sigma_i)$ and cause catastrophic rupture $(\underline{F}_b, \sigma_b)$ in Model 1 specimens.

<u>R</u> i (mm)	<u>R</u> o (mm)	<u>F</u> i (N)	σ _i (N/mm ²) calcula	<u>F</u> b (N) ated	σ_{b} (N/mm ²)	<u>F</u> i (N) meas	<u>F</u> b (N) sured	 (mm)	σ_{b}^{**} (N/mm ²) calc.
0.16	3.0	2.5	0.09	11.9	0.42			1.58	0.65
0.3	3.0	3.0	0.11	11.7	0.41			1.65	0.64
0.435	3.0	3.5	0.12	11.3	0.40	2.9	10.6	1.72	0.63
0.6	3.0	4.0	0.14	11.1	0.39			1.80	0.61
0.78	3.0	4.6	0.16	12.0	0.42	3.9	10.9	1.89	0.60
1.0	3.0	5.1	0.18	11.6	0.41	4.7	11.2	2.00	0.58
1.16	3.0	5.2	0.18	11.5	0.41	5.5	10.6	2.08	0.57
1.5	3.0	5.5	0.19	10.7	0.38			2.25	0.55
2.0	3.0	5.1	0.18	9.1	0.32			2.42	0.53
2.4	3.0	4.2	0.15	8.1	0.29			2.64	0.51
0.435	0.545	0.4	0.43	0.6	0.62			0.49	1.17
0.435	1.45	1.3	0.20	3.7	0.56			0.94	0.85
0.435	2.18	2.2	0.15	6.8	0.46			1.31	0.72
0.435	5.6	6.9	0.07	28.6	0.29	11.3	32.3	3.02	0.47

* : value when <u>G</u> is a minimum.

** : calculated from Eq. 3 at the critical crack radius \underline{a}_c .

<u>R</u> i (mm)	<u>R</u> 0 (mm)	$\frac{F_i}{(N)}$ calculated	<u>F</u> i (N) measured
0.15	3.0	6.0	
1.0	3.0	11.8	
2.0	3.0	16.8	
0.435	2.0	7.5	
0.435	3.0	8.4	9.5
0.435	4.0	9.9	10.3

.

Table 2. Calculated and measured forces \underline{F}_i to initiate a crack in Model 2 specimens.

Figure Legends

- 1 (a) Sketch of the first model. A resin block contains two rods with their ends touching, representing a broken fiber.
 - (b) Sketch of the second model. A rigid tube, representing unbroken fibers, surrounds a resin cylinder containing a broken fiber.
- 2. Fracture energy \underline{G}_{C} vs. tearing rate for silicone rubber samples cured at 110^OC (0) and at room temperature (\bullet).
- 3. Force-displacement relation for a sample with a propagating crack. $\underline{R}_i = 0.435 \text{ mm}; \underline{R}_o = 3.0 \text{ mm}.$
- 4. Variation of residual compressive force with temperature. $\underline{R}_i = 0.435 \text{ mm}; \underline{R}_o = 3.0 \text{ mm}.$
- 5. Sketch of finite element grid for a sample containing a crack of radius <u>a</u>. The shaded area represents the embedded rod (fiber).
- 6. Comparison of FEM results for stress intensity factor \underline{Y} ' (defined by Equation 9) for a penny-shaped crack in a solid cylinder, with values calculated from Equation 10. Dimensions of cylinder: radius = 3.0 mm; length = 50 mm.
- 7. Calculated compliances of Model 1 specimens vs. dimensionless crack radius $(\underline{a-R_i})/(\underline{R_o-R_i})$. Rod radius $\underline{R_i}$ as shown; external radius $\underline{R_o} = 3.0$ mm.
- 8. Cracks photographed at different stages for a sample with rod radius $\underline{R}_i = 0.435$ mm, external radius $\underline{R}_o = 3.0$ mm.

- 9. Comparison of measured and calculated compliances as a function of crack radius, for a sample of 3.0 mm external radius. The vertical line denotes the rod radius \underline{R}_i .
- 10. Calculated energy release rate <u>G</u> for unit applied load vs. dimensionless crack radius $(\underline{a - R_i})/(\underline{R_o - R_i})$. Rod radius $\underline{R_i}$ as shown; sample radius $\underline{R_o} = 3.0$ mm.
- 11. Comparison of applied load \underline{F} vs. crack radius, calculated by FEM, with experimentally measured values. The vertical line denotes the rod radius \underline{R}_i .
- 12. Comparison of applied load \underline{F} vs. crack radius, calculated by FEM, with experimentally measured values. The vertical line denotes the rod radius \underline{R}_i .
- 13. Compliances calculated by FEM for Model 2 specimens, with various values of rod radius \underline{R}_i . External radius $\underline{R}_o = 3.0$ mm.
- 14. Calculated energy release rate <u>G</u> vs. dimensionless crack radius for Model 2 specimens subjected to unit applied load. Rod radius <u>R</u>_i as shown; resin radius <u>R</u>_o = 3.0 mm.
- 15. Calculated and measured values of applied load F vs. crack radius for a Model 2 specimen. Rod radius \underline{R}_i (represented by the vertical line) = 0.435 mm; resin radius \underline{R}_0 = 3.0 mm.
- 16. Sketch of light ray paths 1 and 2 to indicate optical magnification of crack by the resin cylinder acting as a lens.
- 17. Calculated magnification $\underline{y}/\underline{y}_{\alpha}$ of the crack radius.

18. Poissonian contraction of the crack radius. Lateral displacement (contraction) δ' of the crack tip vs. longitudinal displacement (extension) δ of the sample, for unit applied load. $\underline{R}_i = 0.435 \text{ mm}; \underline{R}_0 = 3.0 \text{ mm}.$







FIGURE 1 (b)









.











Photographs of crack propagation

Figure 8









0. e 0 0 0 4 4 0.8 4 0 0 0 4 4 0.6 $(a-R_i)/(R_o-R_i)$ FIGURE 13 4 0 4 0.4 4 0 م م 0 0 0.2 : 0.435 : 1.00 R_i(mm) 2.00 4 0 4 004 0 0.0 0.00 0.04 0.08 0.02 0.06 0.10 (N/mm) esneilqmos

0. R_i(mm) O: 0.435 ●: 1.00 ∆: 2.00 0.8 0.4 0.6 (a-R_i)/(R_o-R_i) 0 \triangleleft 0 4 0 0 4 0.2 4 0 4 0 0.0 0 2 3 (J/m²) C 49

FIGURE 14









magnification order, y/y_0



.

(DYN)

DISTRIBUTION LIST

Dr. R.S. Miller Office of Naval Research Code 432P Arlington, VA 22217 (10 copies)

Dr. J. Pastine Naval Sea Systems Command Code 06R Washington, DC 20362

Dr. Merrill K. King Atlantic Research Corp. 5390 Cherokee Avenue Alexandria, VA 22312

Dr. R.L. Lou Aerojet Strategic Propulsion Co. Bldg. 05025 - Dept 5400 - MS 167 P.O. Box 15699C Sacramenta, CA 95813

Dr. R. Olsen Aerojet Strategic Propulsion Co. Bldg. 05025 - Dept 5400 - MS 167 P.O. Box 15699C Sacramento, CA 95813

Dr. Randy Peters Aerojet Strategic Propulsion Co. Bldg. 05025 - Dept 5400 - MS 167 P.O. Box 15699C Sacramento, CA 95813

Dr. D. Mann U.S. Army Research Office Engineering Division Box 12211 Research Triangle Park, NC 27709-2211 Dr. L.V. Schmidt Office of Naval Technology Code 07CT Arlington, VA 22217

JHU Applied Physics Laboratory ATTN: CPIA (Mr. T.W. Christian) Johns Hopkins Rd. Laurel, MD 20707

Dr. R. McGuire Lawrence Livermore Laboratory University of California Code L-324 Livermore, CA 94550

P.A. Miller 736 Leavenworth Street, #6 San Francisco, CA 94109

Dr. W. Moniz Naval Research Lab. Code 6120 Washington, DC 20375

Dr. K.F. Mueller Naval Surface Weapons Center Code Rll White Oak Silver Spring, MD 20910

Prof. M. Nicol Dept. of Chemistry & Biochemistry University of California Los Angeles, CA 90024

Mr. L. Roslund Naval Surface Weapons Center Code R10C White Oak, Silver Spring, MD 2091C

Dr. David C. Sayles Ballistic Missile Defense Advanced Technology Center P.O. Box 1500 Huntsville, AL 35807

(DYN)

DISTRIBUTION LIST

Mr. R. Geisler ATTN: DY/MS-24 AFRPL Edwards AFB, CA 93523

Naval Air Systems Command ATTN: Mr. Bertram P. Sobers NAVAIR-320G Jefferson Plaza 1, RM 472 Washington, DC 20361

R.B. Steele Aerojet Strategic Propulsion Co. P.O. Box 15699C Sacramento, CA 95813

Mr. M. Stosz Naval Surface Weapons Center Code R10B White Oak Silver Spring, MD 20910

MR. S.F.PALOPALI THIOKOL CORP, ELKTON DIV. P.O.BOX 241 ELKTON MD 21921

Dr. Grant Thompson Morton Thiokol, Inc. Wasatch Division MS 240 P.O. Box 524 Brigham City, UT 84302

Dr. R.S. Valentini United Technologies Chemical Systems P.O. Box 50015 San Jose, CA 95150-0015

Dr. R.F. Walker Chief, Energetic Materials Division DRSMC-LCE (D), B-3022 USA ARDC Dover, NJ 07801 Director * US Army Ballistic Research Lab. ATTN: DRXBR-IBD Aberdeen Proving Ground, MD 21005

Commander US Army Missile Command ATTN: DRSMI-RKL Walter W. Wharton Redstone Arsenal, AL 35898

Dr. Ingo W. May Army Ballistic Research Lab. ARRADCOM Code DRXBR - 1BD Aberdeen Proving Ground, MD 21005

Dr. E. Zimet Office of Naval Technology Code 071 Arlington, VA 22217

Dr. Ronald L. Derr Naval Weapons Center Code 389 China Lake, CA 93555

Lee C. Estabrook, P.E. Morton Thiokol, Inc. P.O. Box 30058 Shreveport, Louisiana 71130

Dr. J.R. West Morton Thiokol, Inc. P.O. Box 30058 Shreveport, Louisiana 7!130

Dr. D.D. Dillehay Morton Thiokol, Inc. Longhorn Division Marshall, TX 75670

G.T. Bowman Atlantic Research Corp. 7511 Wellington Road Gainesville, VA 22065

(DYN)

DISTRIBUTION LIST

R.E. Shenton Atlantic Research Corp. 7511 Wellington Road Gainesville, VA 22065

Mike Barnes Atlantic Research Corp. 7511 Wellington Road Gainesville, VA 22065

Dr. Lionel Dickinson Naval Explosive Ordinance Disposal Tech. Center Code D Indian Head, MD 20340

Prof. J.T. Dickinson Washington State University Dept. of Physics 4 Pullman, WA 99164-2814

M.H. Miles Dept. of Physics Washington State University Pullman, WA 99164-2814

Dr. T.F. Davidson Vice President, Technical Morton Thiokol, Inc. Aerospace Group 3340 Airport Rd. Ogden, UT 84405

Mr. J. Consaga Naval Surface Weapons Center Code R-16 Indian Head, MD 20640

Naval Sea Systems Command ATTN: Mr. Charles M. Christensen NAVSEA-62R2 Crystal Plaza, Bldg. 6, Rm 806 Washington, DC 20362

Mr. R. Beauregard Naval Sea Systems Command SEA 64E Washington, DC 20362 Brian Wheatley Atlantic Research Corp. 7511 Wellington Road Gainesville, VA 22065

Mr. G. Edwards Naval Sea Systems Command Code 62R32 Washington, DC 20362

C. Dickinson Naval Surface Weapons Center White Oak, Code R-13 Silver Spring, MD 20910

Prof. John Deutch MIT Department of Chemistry Cambridge, MA 02139

DAVID A. FLANIGAN, DIR.ADV.TECH MORTON THIOKOL INC AEROSPACE GROUP 2475 WASHINGTON BLVD. OGDEN UT 84401

Dr. L.H. Caveny Air Force Office of Scientific Research Directorate of Aerospace Sciences Bolling Air Force Base Washington, DC 20332

Dr. Donald L. Ball Air Force Office of Scientific Research Directorate of Chemical & Atmospheric Sciences Bolling Air Force Base Washington, DC 20332

(DYN)

• بر

DISTRIBUTION LIST

Dr. Anthony J. Mctuszko Air Force Office of Scientific Research Directorate of Chemical & Atmospheric Sciences Bolling Air Force Base Washington, DC 20332

Dr. Michael Chaykovsky Naval Surface Weapons Center Code R11 White Oak Silver Spring, MD 20910

J.J. Rocchio USA Ballistic Research Lab. Aberdeen Proving Ground, MD 21005-5066

B. Swanson
INC-4 MS C-346
Los Alamos National Laboratory
Los Alamos, New Mexico 87545

Dr. James T. Bryant Naval Weapons Center Code 3205B China Lake, CA 93555

Dr. L. Rothstein Assistant Director Naval Explosives Dev. Engineering Dept. Naval Weapons Station Yorktown, VA 23691

Dr. Henry Webster, III Manager, Chemical Sciences Branch ATTN: Code 5063 Crane, IN 47522

Dr. A.L. Slafkosky Scientific Advisor Commandant of the Marine Corps Code RD-1 Washington, DC 20380 Dr. H.G. Adolph Naval Surface Weapons Center Code Rll White Oak Silver Spring, MD 20910

U.S. Army Research Office Chemical & Biological Sciences Division P.O. Box 12211 Research Triangle Park, NC 27709

Dr. John S. Wilkes, Jr. FJSRL/NC USAF Academy, CO 80840

Dr. H. Rosenwasser AIR-320R Naval Air Systems Command Washington, DC 20361

Dr. Joyce J. Kaufman The Johns Hopkins University Department of Chemistry Baltimore, MD 21218

Dr. A. Nielsen Naval Weapons Center Code 385 China Lake, CA 93555

Ser 432/84/340 Revised January 1985

(DYN)

DISTRIBUTION LIST

K.D. Pae High Pressure Materials Research Lab. Rutgers University P.O. Box 909 08854 Piscataway, NJ Dr. John K. Dienes T-3, B216 Los Alamos National Lab. P.O. Box 1663 Los Alamos, NM 87544 A.N. Gent Institute Polymer Science University of Akron Akron, OH 44325 Dr. D.A. Shockey SRI International 333 Ravenswood Ave. Menlo Park, CA 94025 Dr. R.B. Kruse Morton Thiokol, Inc. Huntsville Division Huntsville, AL 35807-7501 G. Butcher Hercules, Inc. P.O. Box 98 Magna, UT 84044 W. Waesche Atlantic Research Corp. 7511 Wellington Road Gainesville, VA 22065 Dr. R. Bernecker

Naval Surface Weapons Center Code R13 White Oak Silver Spring, MD 20910 Prof. Edward Price Georgia Institute of Tech. School of Aerospace Engineering Atlanta, GA 30332

Prof. R.W. Armstrong University of Maryland Dept. of Mechanical Engineering College Park, MD 20742

Herb Richter Code 385 Naval Weapons Center China Lake, CA 93555

G.A. Zimmerman Aeroject Tactical Systems P.O. Box 13400 Sacramento, CA 95813

Prof. Kenneth Kuo Pennsylvania State University Dept. of Mechanical Engineering University Park, PA 16802

T.L. Boggs Naval Weapons Center Code 3891 China Lake, CA 93555

Ser 432/84/340 Revised January 1985

(DYN)

DISTRIBUTION LIST

Dr. C.S. Coffey Naval Surface Weapons Center Code R13 White Oak Silver Spring, MD 20910

D. Curran SRI International 333 Ravenswood Avenue Menlo Park, CA 94025

E.L. Throckmorton Code SP-2731 Strategic Systems Program Office Crystal Mall #3, RM 1048 Washington, DC 23076

R.G. Rosemeier Brimrose Corporation 7720 Belair Road Baltimore, MD 20742

C. Gotzmer Naval Surface Weapons Center Code R-11 White Oak Silver Spring, MD 20910

G.A. Lo 3251 Hanover Street B204 Lockheed Palo Alto Research Lab Palto Alto, CA 94304

R.A.SCHAPERY DEPT. OF AEROSPACE ENGNG. UNIVERSITY OF TEXAS AUSTIN TX 78712

Dr. Y. Gupta Washington State University Department of Physics Pullman, WA 99163 J.M. Culver Strategic Systems Projects Office SSPO/SP-2731 Crystal Mall #3, RM 1048 Washington, DC 20376

Prof. G.D. Duvall Washington State University Department of Physics Pullman, WA 99163

DR.G. LINDSAY NAVAL WEAPONS CENTER CODE 3858 CHINA LAKE CA 93555

Dr. M. Farber 135 W. Maple Avenue Monnovia, CA 91016

W.L.ELBAN DEPT. OF ELECTRICAL ENGNG. LOYOLA COLLEGE 4501 N. CHARLES STREET BALTIMORE ND 21210

Defense Technical Information Center Bldg. 5, Cameron Station Alexandria, VA 22314 (12 copies)

Dr. Robert Polvani National Bureau of Standards Metallurgy Division Washington, D.C. 20234

Director Naval Research Laboratory Attn: Code 2627 Washington, DC 20375 (6 copies)

MR. B. CONLEY ONR RESIDENT REPRESENTATIVE OHIO STATE UNIV. RES. FOUND. 1314 KINNEAR ROAD COLUMBUS, OHIO 43212