

EVALUATION OF ENERGY RELEASE RATE FOR MODE I CRACK PROPAGATION IN GLASS/EPOXY COMPOSITE

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Abstract

Composite resources are replacing regular engineering metals along with alloys for several applications. Their superior specific strength and stiffness characteristics have made them very competitive in the aerospace industry. The primary limitation of fiber reinforced composites is fracture toughness, specifically delamination. Delamination failures are common due to the nature of composite construction. A variety of manufacturing techniques are available to make composites. Generally, all these methods employ a layered stacking of fibers in a primary plane. The interface between these layers is typically not reinforced with fibers and is the source of delamination or interlaminar fracture. Porosity and Other manufacturing related defects also introduce nucleation sites for delamination. Methods exist to evaluate and quantify inter-laminar fracture toughness, both experimentally and analytically. The material property that best represents resistance to delamination is the strain energy release rate (G_C). This can be experimentally obtained and numerically predicted with some success. The primary focus of this study was to characterize and address interlaminar fracture in composites. An interlaminar fracture for mode I crack propagation was analysed for GFRP materials having 0° orientation of fibres. Experimental investigation of G_C was done for 40%, 50% and 60% of fibre volume. Test was conducted using Double Cantilever Beam specimen. Results obtained were again correlated with Numerical analysis using VCCT method. Results of both experimental and numerical methods were in good agreement.

Keywords: S- Glass fiber, Epoxy Resin, Mechanical properties, Hand lay-up.

1. INTRODUCTION

Composites materials are the combination or mixture of two or more dissimilar materials. That are combined in the macroscopic level but these materials chemically not combined together.

The main aim to develop the composite materials in order to reduce the weight of the materials and improve the mechanical and thermal properties of the components and the structural and the structural applications Composite materials are the combinations of reinforcing, fillers, elements, and matrix binder, differing in form on a macro scale. It should be a strong interface bond between the reinforcement and the matrix is obviously desirable, so the matrix must be capable of developing a mechanical or chemical bond with the reinforcement. The reinforcement and the matrix materials should also be chemically compatible.

In these materials one is matrix material and another is reinforcing material. These depend on the applications and material composition will be changed.. Based on this matrix composite materials are further classified in to metal matrix composites, polymer matrix composites and ceramics.

The structure of polymers consists of long molecules with a backbone of carbon atoms linked by covalent bonds. In non-crystalline or amorphous polymers the molecular chains have an entirely random orientation and are cross-linked

occasionally by a few strong covalent bonds and numerous but weaker van der Waals bonds. These weaker bonds break as the temperature reaches a value known as the glass transition temperature, are oriented along preferred directions.

Double Cantilever Beam Specimen

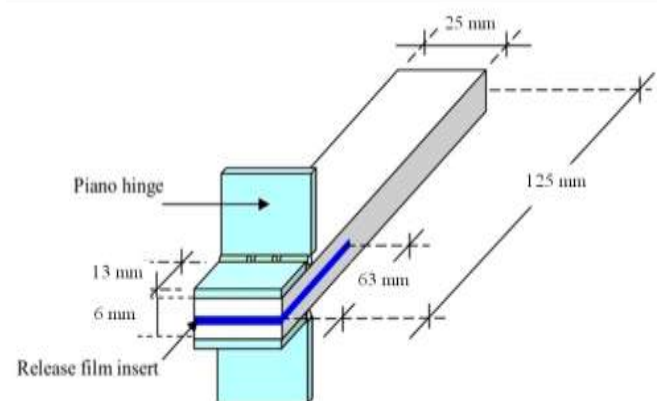


Fig1 DCB specimen

Dimensions of DCB specimen ASTM 5528

Were;

U=total elastic energy in the test specimen,

b= width, and

a= length,

l=125mm
 L=63mm
 a₀=13mm
 h=3mm
 b=25mm

The maximum load anticipated during a DCB test of a material with a known modulus, E₁₁, and anticipated value of G_{IC}, may be estimated by

$$P_{max} = \frac{b}{a} \sqrt{\frac{E_{11} G_{IC} h^3}{96}}$$

Total volume of the specimen V=12500mm³=12.5cm³

Density of S glass ρ_g=2.45 (g/cm³)

Mass of S glass required to prepare specimen with volume fraction, v_f= v_fX15X2.45=36.75v_fg

Mass of S glass used in for fiber volume fraction

Volume fraction V_f = 0.4 E₁₁ = 36.

Volume fraction V_f = 0.4 E₁₁ = 36.2

V_f = 0.4 5 E₁₁ = 40.35

V_f = 0.5 E₁₁ = 44.5

V_f = 0.6 E₁₁ = 52.8

Properties of Epoxy Resin

Density ρ_m=1.2 (g/cm³)

Young's Modulus E_m= 3 (Gpa)

Poisson's Ratio ν_m = 0.38

Shear strength - τ_m=0.034 (Gpa)

Tensile Strength - σ_{tm} = 0.072 (Gpa)

Axial Compressive strength σ_{mc} = 1.550 (Gpa)

Properties of S-Glass Fiber

Density ρ_g=2.45 (g/cm³)

Young's Modulus E_f = 86 (Gpa)

Poisson's Ratio ν_f = 0.22

Shear strength - τ_f=0.035 (Gpa)

Axial Compressive strength σ_{fc} = 1.550 (Gpa)

From the properties of S glass and epoxy, based on the their volume fraction elastic moduli are obtained.

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a. Longitudinal young's modulus, E₁=E_fV_f+E_mV_m

b. Major Poisson's ratio,

$$\nu_{12} = \nu_f V_f + \nu_m V_m$$

The study was performed for 0° orientation of glass fiber having 9 elastic constants in numerical method.

$$E_{11} = V_f E_f + V_m E_m$$

$$E_{22} = E_{33} = \frac{E_{f22} E_m}{E_{f22} V_m + E_m V_f}$$

$$G_{12} = G_{13} = \frac{G_{f12} G_m}{G_{f12} V_m + G_m V_f}$$

$$G_{23} = \frac{G_{f23} G_m}{G_{f12} V_m + G_m V_f}$$

$$\nu_{12} = \nu_{13} = V_f \nu_{f12} + V_m \nu_m$$

$$\nu_{23} = V_f \nu_{f23} + V_m \nu_m$$

2. FABRICATION AND TESTING

1. The fabrication of the polymer matrix combined was prepared at extent temperature.
2. Prepare the mould cavity according to the specimen dimensions.
3. Fibers are chopped in uni directional and placed on the mould cavity.
4. The epoxy and resins mixed together and applied on the fibers in layer by layer.
5. PTFE is used as the releasing agent, load is kept on the cavity for he certain interval of time.

Testing Procedure

1. Measure the width and thickness of each specimen to the nearest at the midpoint and from either end.
2. Coat both edges of the specimen just ahead to identify the crack length..
3. The specimen is fixed in between the two hinges , aligned center to the specimen.
4. Once load gets increased the crack starts propagating gradually. Reading should be note down.
5. As the load increases the specimen breaks at that point.



Fig 2 UTM machine

Machine Make: Instron
Model:5569
Capacity:50KN
Universal Testing Machine

3. RESULTS AND DISCUSSIONS

Numerical results and Experimental results are compared to plot the graph. Numerical method is done by Virtual Crack Closure Technique and compared both results.

3.1 Numerical Analysis of Double Cantilever Beam for different volume fraction of Fiber

3.1.1 Fiber Volume Fraction $V_f=0.4$

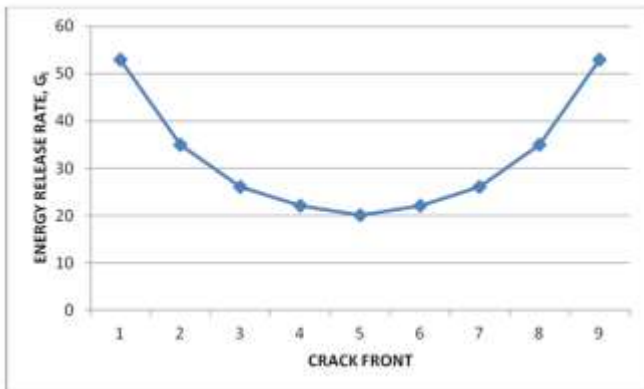
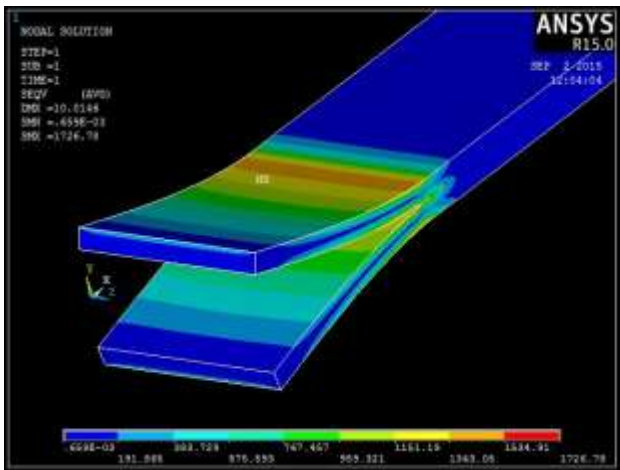


Fig 3: Graph gives the pattern of G value along the crack front. Around 51.2 J/m².

3.1.2 Fiber Volume Fraction $V_f=0.45$

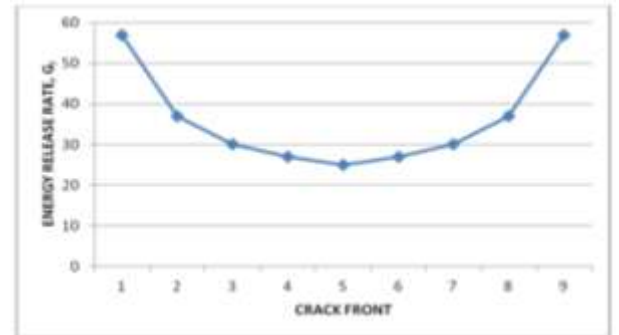
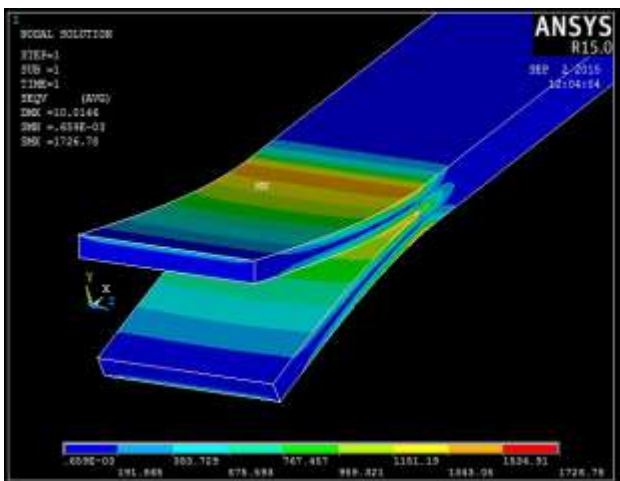


Fig 4: Graph gives the pattern of G value along the crack front. Around 56.6 J/m².

3.1.3 Fiber Volume Fraction $V_f=0.5$

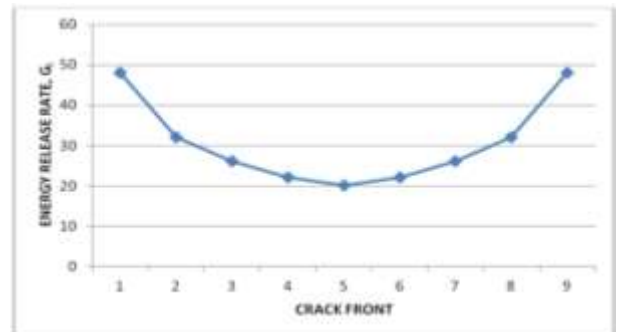
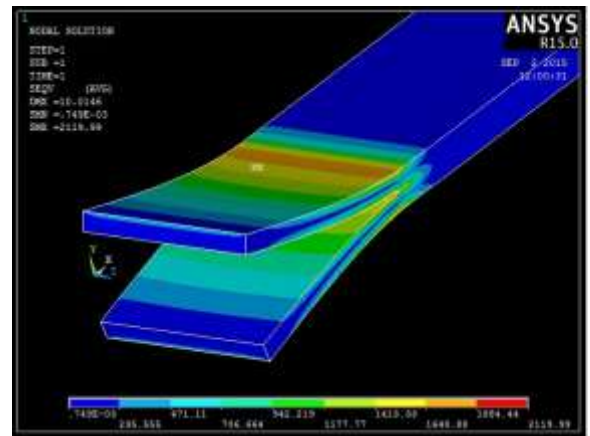
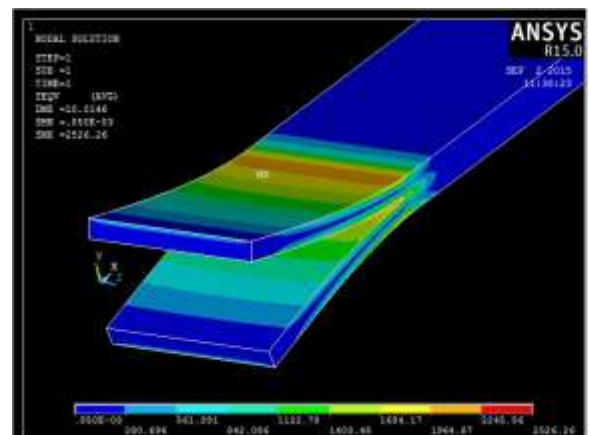


Fig 5: Graph gives the pattern of G value along the crack front. Around 47.4 J/m².

3.1.4 Fiber Volume Fraction $V_f=0.6$



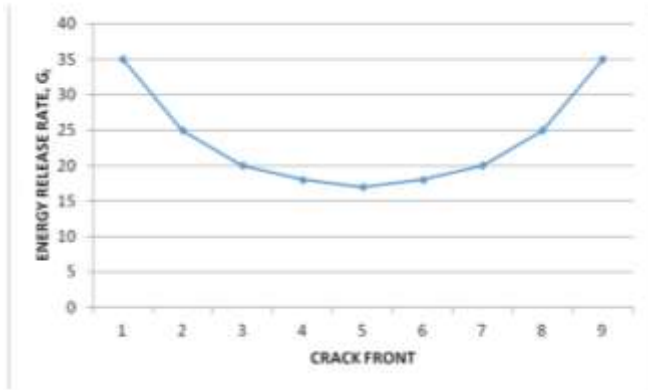


Fig 6: Graph gives the pattern of G value along the crack front. Around 35.1 J/m².

3.2.3 Fiber Volume Fraction V_f=0.5

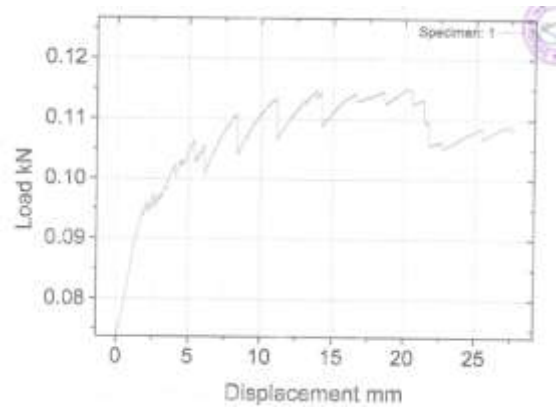


Fig 9

$$G_{1c} = 48.71 \text{ J/m}^2$$

3.2 Experimental Results Calculated using above Mentioned Formulas

3.2.1 Fiber Volume Fraction V_f=0.4

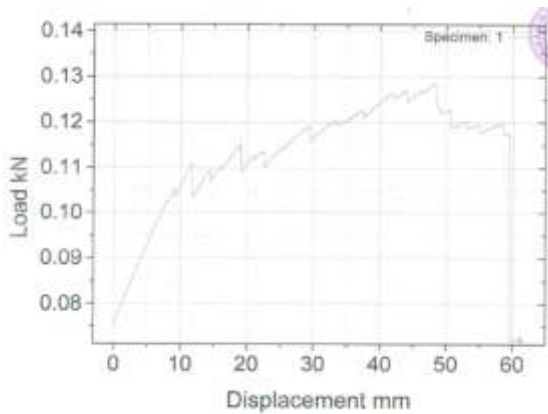


Fig 7

$$G_{1c} = 96 \frac{P_{max} a^2}{E_{11} h^3 b^2} G_{1c} = 53.64 \text{ J/m}^2$$

3.2.4 Fiber Volume Fraction V_f=0.6

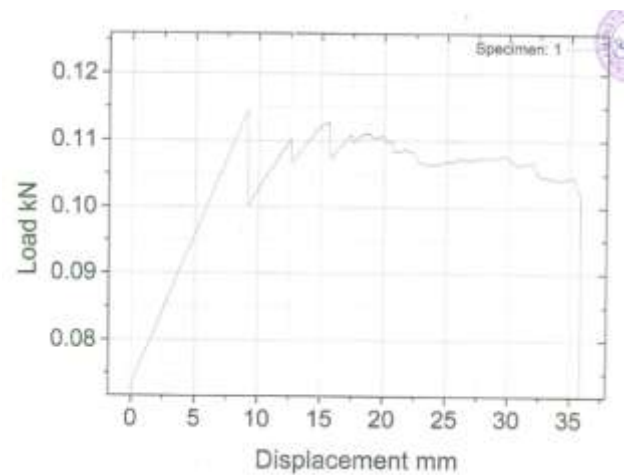


Fig 10

$$G_{1c} = 34.21 \text{ J/m}^2$$

3.2.2 Fiber Volume Fraction V_f=0.45

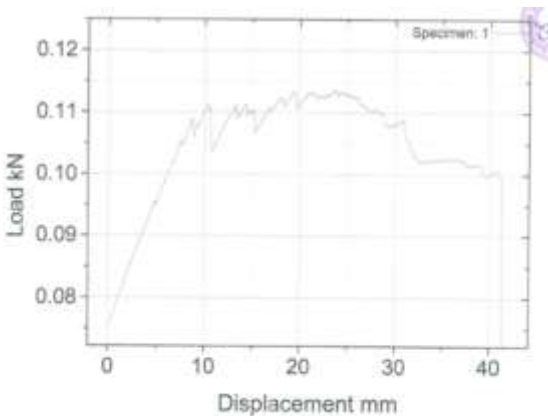


Fig 8

$$G_{1c} = 57.07 \text{ J/m}^2$$

3.3 Comparison of Numerical Results with Experimental Results

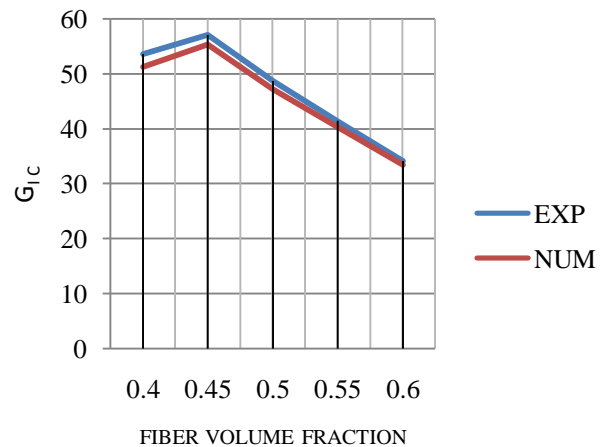


Fig 11 comparison graph

4. CONCLUSION

1. An wide-ranging analysis for Mode I has been performed in arrange to revise the model of Energy Release rate deviation with admiration to modify in fiber volume throughout numerical method and also for the same fiber volume fraction experimental methodology is done.
2. The revise was performed for 0° orientation of glass fiber having 9 elastic constants in numerical method.
3. The fibers are hand lapped and specimen is prepared for experimental method. It clear shows at 45 % fiber the energy release rate will be more both in numerical and experimental in mode I.
4. The results obtained so far are quite satisfactory. The experimental and numerical results were compared and found almost similar.
5. Hence it is clear from both numerical and experimental as percentage of fiber increase Energy release rate decreases in Mode I.

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