

# Fracture Behavior of 2124Al-10vol% SiCp Composite Tensile Specimens for Different Heat Treatment Conditions

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## Abstract

*This article presents experimental details on 2124Al-10vol% SiCp composite tensile specimens, which were made by squeeze casting process. Cast metal composites are generally very brittle and have poor mechanical properties. To strengthen the matrix and improve ductility, the specimens were heat-treated by solutionizing and aging. Experiments were carried out at 3 different solutionizing temperatures, solutionizing time, aging temperature and aging time to examine the influence of heat treatment process parameters on the notched and unnotched tensile strength of the composite. Compact tension (CT) specimens with different crack lengths, crack mouth widths and thicknesses were tested to evaluate the fracture toughness of the composite. Experiments were planned according to the Taguchi's L9 OA (orthogonal array). Factographs of tensile and CT specimens were made after the test using a scanning electron microscope (SEM). Energy dispersive X-ray analysis was carried out at different locations of the fractured specimens. The EDAX spectrum shows the chemical composition of inclusions and matrix components.*

**Keywords:** EDAM spectrum; Fracture toughness; Metal matrix composites; SEM; Taguchi method; Tensile strength.

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## INTRODUCTION

Silicon carbide (SiCp) reinforced aluminum matrix composites have high strength, high specific modulus, high wear resistance, and excellent corrosion resistance, which have broad application prospects in automotive, machinery, aviation, and electronic packaging fields [1, 2]. Addition of SiC particles to aluminum composites improves hardness, density, and compressive strength [3–6]. The mechanical properties of SiC particle-reinforced metal matrix composites (MMCs) are influenced by microstructure and interfacial reactions [7]. The production techniques of centrifugal casting, powder metallurgy, and stir casting offer effective ways to produce aluminum SiC particle MMCs with enhanced properties [8–11]. Aluminum SiC particle MMCs have potential applications in the aerospace industry due to their high strength, lightweight design, wear resistance, and corrosion resistance [12–14].

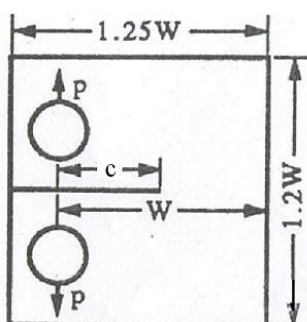
Al-SiC 2124 composites are usually made by the powder metallurgy route, avoiding the use of liquid metal, which would create a brittle reaction zone of  $Al_4C_3$  around the reinforcement. Prasad and Sasidhara [15] have compiled flow stress data for 2124 aluminum with varying volume fractions of SiC particulate (15  $\mu\text{m}$ ) reinforcements in the vacuum hot pressed and extruded at  $500^\circ\text{C}$  for 30 minutes. The nominal composition of the matrix (2124Al) in wt% was 4.2 Cu, 0.99 Mg, 0.72Mn, 0.16Fe, 0.13 Si, 0.03Zn, balance aluminum. 2124AA with 10vol% SiCp exhibits large, grained matrix structure, which restricts the occurrence of super plasticity due to dynamic recrystallization. The material exhibits flow instability in the temperature range  $340\text{--}420^\circ\text{C}$  and at strain rates higher than  $1\text{s}^{-1}$ . 2124 Al-30vol%SiCp metal matrix composite is difficult to process due to its high SiCp content. Murty et al. [16] investigated the hot working characteristics of 2124Al-SiCp metal matrix composites. The mechanical properties of AlMMCs generally lie somewhere between those of unreinforced aluminum and titanium alloys. However, it is possible to alter the balance of properties by careful selection of matrix alloy and level of reinforcement.

Inspired by the work of previous researchers, this paper investigates the fracture behavior of 2124Al-10vol% SiCp composite tensile specimens for different heat treatment conditions. To strengthen the matrix and improve ductility, the specimens were heat-treated by solutionizing and aging. Experiments were conducted to verify the influence of heat treatment process parameters on the notched and unnotched tensile strength of the composites and presented the SEM factographs of the tested samples. Energy dispersive X-ray analysis was performed on fractured specimens. The EDAX spectrum shows the chemical composition of inclusions and matrix components.

### SPECIMEN TESTING OF 2124Al-10vol% SiCp COMPOSITES

Compact tension (CT) specimens (see Figure 1) with different crack lengths, crack mouth widths and thicknesses were tested to evaluate the fracture toughness of the composite. Tests were planned according to the Taguchi's  $L_9$  OA (orthogonal array). Tensile strength and fracture toughness properties of the 2124Al-10vol% SiCp composite were generated for different heat-treatment conditions. The samples were heat-treated by solutionizing and aging using an Indfur electric furnace. Tensile specimens were tested on a 50T capacity UTM. Fracture toughness testing was performed on an Instron Servo hydraulic controlled dynamic testing machine (8801 model).

Nine blanks of  $50 \times 50 \times 7.5$  mm,  $50 \times 50 \times 10$  mm,  $50 \times 50 \times 12$  mm were cut from the extrusion and machined to final dimensions of CT specimens as per ASTM E399 standards. Specimens were notched to L-T orientation. Special care was taken when pre-cracking the specimens with fatigue cracks of controlled lengths. Because early attempts to produce pre-crack by conventional methods using cyclic tensile loading failed due to the rapid propagation of fatigue cracks. To control this pre-cracking problem, fatigue loading with both minimum and maximum loads in compression was used first to initiate pre-cracks. After that, the initial pre-cracking will stop by self-arrest. Afterwards, the pre-crack can be safely extended to the desired length by additional fatigue loading in tension. Following the above procedure, the specimens were successfully pre-cracked in compression with cyclic loadings of 920 and 5.4 kgf. When the crack self-arrested, the loads were changed to 261 and 27.6 kgf in tension to complete the pre-cracking process. In all cases, the cyclic frequency was 6 Hz.



**Figure 1.** Compact Tension (CT) Specimen.

## RESULTS AND DISCUSSION

In the plan of experiments, four factors (viz., solutionizing temperature, solutionizing time, aging temperature and aging time) were selected at three levels. Table 1 gives the factors and the assignment of the corresponding level for which the values taken by the factors. Before selecting an OA (orthogonal array), the minimum number of experiments to be conducted shall be fixed, which is given by [17]

$$N_{\text{Taguchi}} = 1 + \text{Number of factors} \times (\text{number of levels} - 1) = 1 + 4 \times (3-1) = 9.$$

The OA chosen was the  $L_9(3^4)$ . The plane of experiments in Table 2 is made of 9 tests (array rows) where the first column was assigned to solutionizing temperature, the second to the solutionizing time, third to aging temperature and fourth to aging time.

The stress intensity factor ( $K_I$ ) for Compact Tension (CT) specimen is [18]

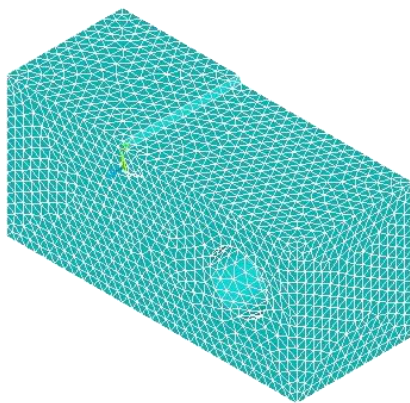
$$K_I = \frac{P}{B\sqrt{W}} Y\left(\frac{c}{W}\right), \quad (1)$$

where  $Y(\xi) = (2 + \xi)(1 - \xi)^{-\frac{3}{2}}(0.886 + 4.64\xi - 13.32\xi^2 + 14.72\xi^3 - 5.6\xi^4)$ .  $P$  is the load;  $B$  is the thickness;  $W$  is the width and  $c$  is crack length of the CT specimen. Using ANSYS, finite element analysis (FEA) has been carried out to evaluate  $K_I$  of the CT specimen. Half the model (see Figure 2) is considered and applied the symmetry boundary conditions. FEA results are found to be in good agreement with the expression (1) for  $K_I$ .

The elastic nominal (net section) stress at failure,  $\sigma_{nf}$ , for CT specimen is [19–22]:

$$\sigma_{nf} = \frac{2P_{max}}{BW} \left(2 + \frac{c}{W}\right) \left(1 - \frac{c}{W}\right)^{-2} \quad (2)$$

where  $P_{max}$  is the failure load. The fracture toughness ( $K_Q$ ) is obtained by substituting the failure load,  $P_{max}$  for  $P$  in equation (1).



**Figure 2.** Half the FE model of CT specimen.

**Table 1.** Assignment of the levels to the factors.

Level	Factors			
	Solutionizing		Aging	
	Temperature (°C)	Time (hrs)	Temperature (°C)	Time (hrs)
1	500	6	180	10
2	520	8	200	12
3	540	10	220	14

Table 2 gives tensile strength ( $\sigma_0$ ), failure load ( $P_{max}$ ) of CT specimens, and fracture toughness ( $K_Q$ ) of the composite. Using the measured output responses from Table 2, analysis of variance (ANOVA) is performed and presented the results in Table 3. Solutionized temperature ( $S_T$ ) has maximum influence on  $\sigma_0$  with 80 % Contribution, while 65 %Contribution on  $K_Q$ . The influence of other parameters ( $S_t$ ,  $A_T$  and  $A_t$ ) on  $\sigma_0$  are: 12.5%, 2 % and 5.5 % respectively. The influence of parameters ( $S_t$ ,  $A_T$  and  $A_t$ ) on  $K_Q$  are: 17.4 %, 2.7 % and 15.4 % respectively. The grand mean value of  $\sigma_0= 265.4$  MPa and the grand mean value of  $K_Q= 18.06 \text{ MPa}\sqrt{\text{m}}$ . Figure 3 shows the fractured CT specimen.

From the mean values of the output responses ( $\sigma_0$ and  $K_Q$ ) in ANOVA Table 3, it is possible to estimate the output responses for the specified levels of the process parameters using the additive law [17]:

$$\hat{\varphi} = \sum_{i=1}^{n_p} \bar{\varphi}_{ik} - (n_p - 1)\varphi_g \quad (3)$$

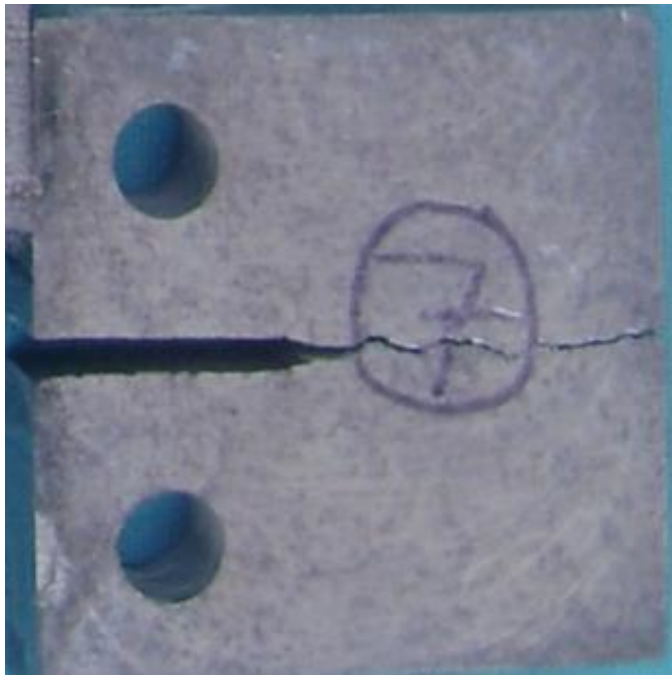
Here,  $\hat{\varphi}$  is the estimate of the output response.  $\varphi_g$ is the grand mean of the output response.  $\bar{\varphi}_{ik}$ is the mean value of the output response corresponding to  $i^{\text{th}}$  process parameter (i.e.,  $i = 1$  for  $S_T$ ;  $i = 2$  for  $S_t$ ;  $i = 3$  for  $A_T$ ; and  $i = 4$  for  $A_t$ ) and  $k^{\text{th}}$  level (i.e.,  $k = 1,2,3$ ). Estimates of  $\sigma_0$ and  $K_Q$  using equation (3) are exactly matching with the measured data in Table 2.

**Table 2.** Taguchi's  $L_9$  OA and values of tensile strength ( $\sigma_0$ ), failure load ( $P_{max}$ ) and fracture toughness ( $K_Q$ ) of CT specimen (Width (W) = 32 mm; Thickness (B) = 7.5 mm; Initial crack size (c) = 10 mm).

Test runs	Solutionizing		Aging		Tensile strength, $\sigma_0$ (MPa)	Failure load, $P_{max}$ (kN)	Fracture toughness, $K_Q$ ( $\text{MPa}\sqrt{\text{m}}$ )
	Temp., $S_T$ ( $^{\circ}\text{C}$ )	Time, $S_t$ (hrs)	Temp., $A_T$ ( $^{\circ}\text{C}$ )	Time, $A_t$ (hrs)			
1	500	6	180	10	277	4.294	18.58
2	500	8	200	12	335	3.338	14.44
3	500	10	220	14	300	3.169	13.71
4	520	6	200	14	177	3.784	16.37
5	520	8	220	10	222	4.065	17.59
6	520	10	180	12	209	3.812	16.49
7	540	6	220	12	298	5.800	25.10
8	540	8	180	14	314	4.309	18.65
9	540	10	200	10	257	5.001	21.64

**Table 3.** ANOVA results on performance characteristics ( $\sigma_0$  and  $K_Q$ )

Parameters	1 <sup>st</sup> Mean	2 <sup>nd</sup> Mean	3 <sup>rd</sup> Mean	SoS	% Contribution
<b>Tensile strength, <math>\sigma_0</math> : Grand mean = 265.4 MPa</b>					
$S_T$ ( $^{\circ}\text{C}$ )	304.0	202.7	289.7	18043	80.0
$S_t$ (hrs)	250.7	290.3	255.3	2820	12.5
$A_T$ ( $^{\circ}\text{C}$ )	266.7	256.3	273.3	440	2.0
$A_t$ (hrs)	252.0	280.7	263.7	1247	5.5
<b>Fracture toughness, <math>K_Q</math> : Grand mean = 18.06 <math>\text{MPa}\sqrt{\text{m}}</math></b>					
$S_T$ ( $^{\circ}\text{C}$ )	15.577	16.817	21.797	65.0	65.0
$S_t$ (hrs)	20.017	16.893	17.280	17.4	17.4
$A_T$ ( $^{\circ}\text{C}$ )	17.907	17.483	18.800	2.7	2.7
$A_t$ (hrs)	19.270	18.677	16.243	15.4	15.4



**Figure 3.** Fractured CT specimen.

Considering the mean values of  $\sigma_0$  and  $K_Q$  from ANOVA Table 3 and the additive law in equation (3), empirical relationships are developed in terms of  $S_T$ ,  $S_t$ ,  $A_T$  and  $A_t$  in the form:

$$\sigma_0 = 233.67 - 7.1667\xi_1 + 94.1667\xi_1^2 + 2.3333\xi_2 - 37.3333\xi_2^2 + 3.3333\xi_3 + 13.6667\xi_3^2 + 5.8333\xi_4 - 22.8333\xi_4^2 \quad (4)$$

$$K_Q = 15.68 + 3.11\xi_1 + 1.87\xi_1^2 - 1.3683\xi_2 + 1.7555\xi_2^2 + 0.4467\xi_3 + 0.87\xi_3^2 - 1.5133\xi_4 - 0.92\xi_4^2 \quad (5)$$

Here,  $\xi_1 = 0.05S_T - 26$ ;  $\xi_2 = 0.5S_t - 4$ ;  $\xi_3 = 0.05A_T - 10$ ; and  $\xi_4 = 0.5A_t - 6$ .

From the mean values of the output responses in ANOVA Table-3, maximum  $\sigma_0$  mean values correspond to the 1<sup>st</sup> level of  $S_T$ , 2<sup>nd</sup> level of  $S_t$ , 3<sup>rd</sup> level of  $A_T$  and 2<sup>nd</sup> level of  $A_t$ . Hence,  $\sigma_{0\max}$  (maximum  $\sigma_0$ ) can be achieved for a set of parameters ( $S_{T1}S_{t2}A_{T3}A_{t2}$ ). Number subscripts indicate the level of the parameters. Similarly, a set of parameters ( $S_{T3}S_{t1}A_{T3}A_{t1}$ ) was found for maximum  $K_Q$  ( $K_{Q\max}$ ). Two different sets of parameters found for  $\sigma_{0\max}$  and  $K_{Q\max}$ . The designer needs a set of parameters to achieve simultaneously  $\sigma_{0\max}$  and  $K_{Q\max}$ . Hence, multi-objective optimization scheme [23–26] is appropriate to have a set of parameters for achieving  $\sigma_{0\max}$  and  $K_{Q\max}$ . These objectives have different units of measurement. To handle such problems, all the output responses are normalized and converted to maximization of a single objective function [25]:

$$\zeta = \omega_1\zeta_1 + \omega_2\zeta_2 \quad (6)$$

Here,  $\zeta_1 = \frac{\sigma_0}{\sigma_{0\max}}$ ;  $\zeta_2 = \frac{K_Q}{K_{Q\max}}$ ;  $\omega_1$  and  $\omega_2$  are the positive weighing factors such that  $\omega_1 + \omega_2 = 1$ ;  $\sigma_{0\max} = 352$  MPa; and  $K_{Q\max} = 25.69$  MPa $\sqrt{m}$ . Maximizing  $\zeta$ , results in high  $\sigma_0$  and high  $K_Q$ . Assuming  $\omega_1 = \omega_2 = 0.5$ , and using mean values of  $\sigma_0$  and  $K_Q$  from ANOVA Table 3 in equation (6), ANOVA results for  $\zeta$  are obtained (see Table 4). A set of parameters ( $S_{T3}S_{t1}A_{T3}A_{t2}$ ) found from the maximum mean values of  $\zeta$ . Hence, optimal parameters to attain maximum  $\zeta$  are:  $S_T = 540^\circ\text{C}$ ;  $S_t = 6$  hours;  $A_T = 220^\circ\text{C}$ ; and  $A_t = 12$  hours. The output responses corresponding to the set of optimal parameters obtained from equations (4) and (5) are:  $\sigma_0 = 297.84$  MPa and  $K_Q = 25.1$  MPa $\sqrt{m}$  (see Table 5), which are matching well with the measured data.

**Table 4.** ANOVA for  $\zeta$ 

Parameters	1 <sup>st</sup> Mean	2 <sup>nd</sup> Mean	3 <sup>rd</sup> Mean
S <sub>T</sub> (°C)	0.7350	0.6152	0.8357
S <sub>t</sub> (hrs)	0.7456	0.7412	0.6990
A <sub>T</sub> (°C)	0.7273	0.7044	0.7542
A <sub>t</sub> (hrs)	0.7330	0.7622	0.6907

**Table 5.** Estimates of strength ( $\sigma_0$ ) and fracture toughness ( $K_Q$ )

Solutionizing		Aging		Strength	Fracture toughness
Temp., S <sub>T</sub> (°C)	Time, S <sub>t</sub> (hrs)	Temp., A <sub>T</sub> (°C)	Time, A <sub>t</sub> (hrs)	$\sigma_0$ (MPa)	$K_Q$ (MPa $\sqrt{m}$ )
<i>Case study-I: Single-objective optimization</i> Set of optimal parameters (S <sub>T1</sub> S <sub>t2</sub> A <sub>T3</sub> A <sub>t2</sub> ) for maximum $\sigma_0$					
500	8	220	12	351.84	15.76
<i>Case study-II: Single-objective optimization</i> Set of optimal parameters (S <sub>T3</sub> S <sub>t1</sub> A <sub>T3</sub> A <sub>t1</sub> ) for maximum $K_Q$					
540	6	220	10	269.17	25.69
<i>Case study-III: Multi-objective optimization</i> Set of optimal parameters (S <sub>T3</sub> S <sub>t1</sub> A <sub>T3</sub> A <sub>t2</sub> ) for maximum $\sigma_0$ and maximum $K_Q$					
540	6	220	12	297.84 (298) <sup>+</sup>	25.1 (25.1)
* Measured data					

**Table 6.** Effect of thickness (B) on the fracture toughness of the CT specimens for S<sub>T</sub> = 500°C; S<sub>t</sub> = 12 hours; A<sub>T</sub> = 200°C; and A<sub>t</sub> = 12 hours. (Width, W = 32 mm; Tensile strength,  $\sigma_0$  = 282 MPa)

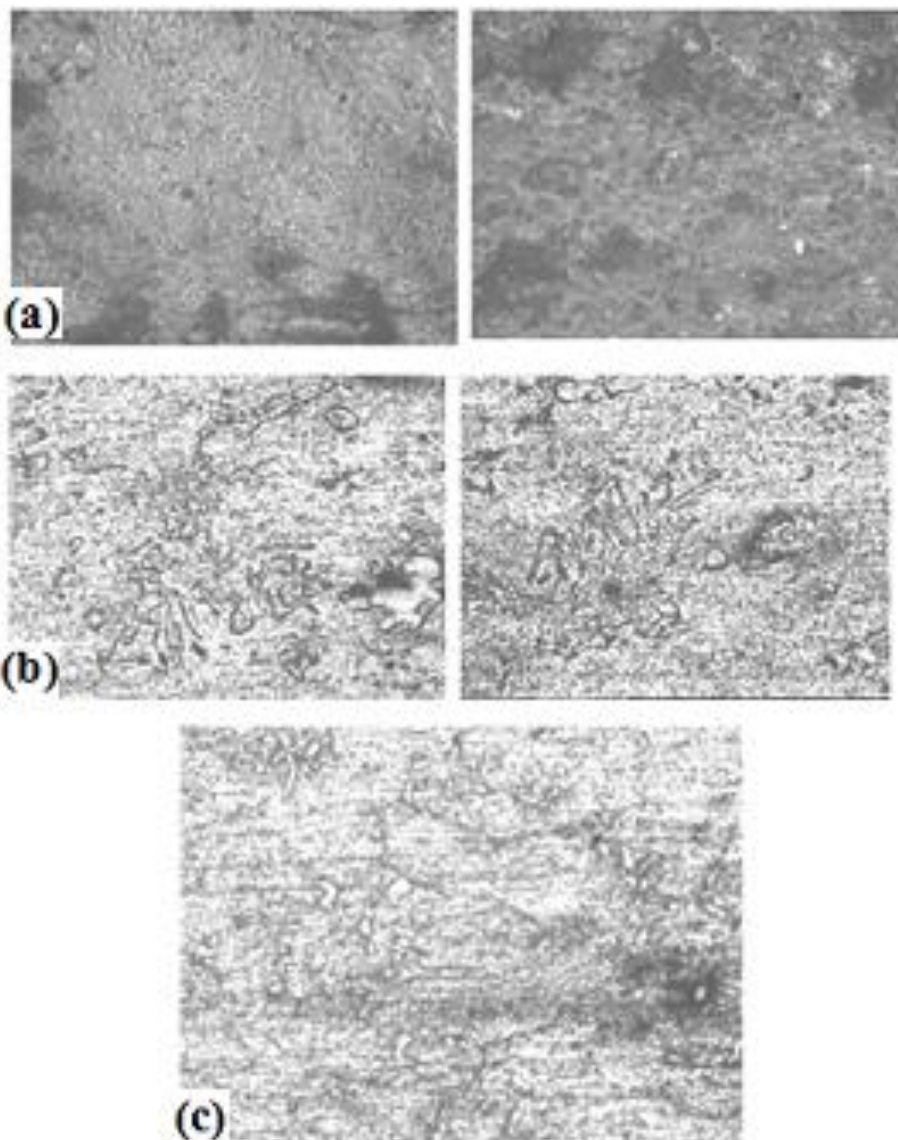
Crack Length, c (mm)	Thickness, B (mm)	Crack Mouth Width (mm)	Failure load, P <sub>max</sub> (kN)	Net section Stress, $\sigma_{nf}$ (MPa)	Fracture toughness, $K_Q$ (MPa $\sqrt{m}$ )
10	7.5	1.2	4.953	201.9	21.43
13.33	10	1.4	5.030	223.2	21.39
16	12	1.6	4.990	259.9	22.45
11	7.5	1.4	4.956	224.8	23.24
14.66	10	1.6	4.325	226.3	20.63
17.6	12	1.2	4.580	300.4	24.25
12	7.5	1.6	4.241	214.9	21.55
16	10	1.2	3.946	246.6	21.31
19.2	12	1.4	3.697	312.9	23.52

Tensile strength ( $\sigma_0$ ) of the un-heat-treated composite is 163 MPa, while it is 282 MPa for the heat-treatment process: S<sub>T</sub> = 500°C; S<sub>t</sub> = 12 hours; A<sub>T</sub> = 200°C; and A<sub>t</sub> = 12 hours. The effect of thickness on the fracture toughness ( $K_Q$ ) of the CT specimens is shown in Table 6. It is evidenced from FEA results that the crack mouth width has no effect on  $K_I$ . P<sub>max</sub> decreases with increasing crack size and the average  $K_Q = 22.2 \text{ MPa}\sqrt{m}$ .

CT specimens were subjected to hot pressing. Table 7 gives  $K_Q$  values for different parameters (viz., temperature and % of compression). The dimensions of the specimens are: Width (W) = 32 mm; Thickness (B) = 7.5 mm; Crack mouth width = 1.2 mm; and Initial crack size (c) = 10 mm. The maximum  $K_Q = 24.35 \text{ MPa}\sqrt{m}$  for the hot-pressing parameters: Temperature = 500°C and 50% of compression.

**Table 7.** Effect of hot pressing on the fracture toughness ( $K_{Ic}$ ) of the CT specimens.

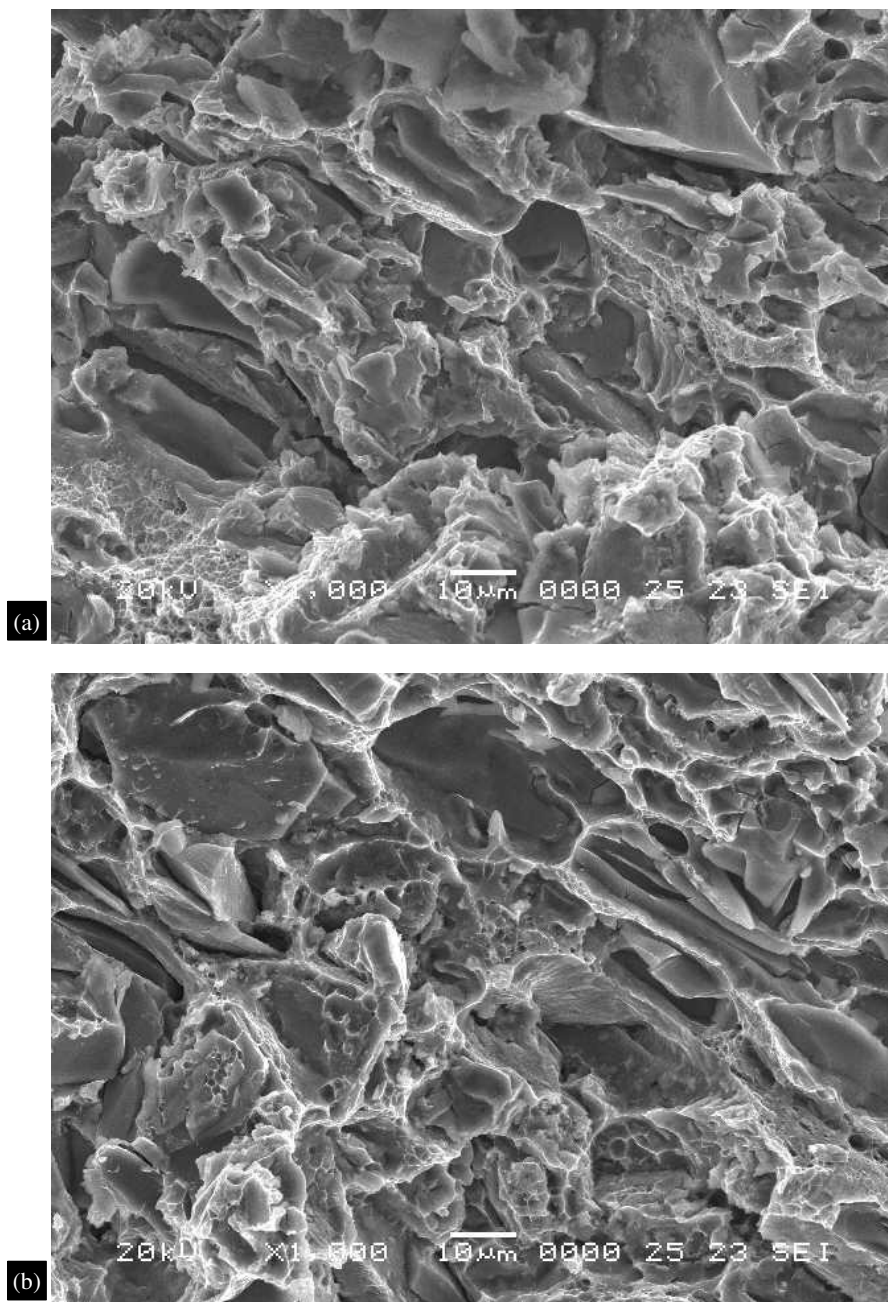
Hot pressing parameters		Failure load, $P_{max}$ (kN)	Net section Stress, $\sigma_{nf}$ (MPa)	Fracture toughness, $K_{Ic}$ (MPa $\sqrt{m}$ )
Temperature (°C)	% of compression			
450	15	2.547	103.8	11.02
450	25	4.920	200.6	21.29
500	25	4.726	192.9	20.45
500	50	5.628	229.5	24.35
550	15	5.236	213.5	22.66
550	25	3.717	151.5	16.08
550	50	5.194	211.8	22.47



**Figure 4.** Optical microstructures at different heat-treatment processes: (a) un-heat-treated; (b)  $S_T = 500^\circ\text{C}$ ;  $S_t = 8$  Hrs;  $A_T = 200^\circ\text{C}$ ;  $A_t = 12$  Hrs; and (c)  $S_T = 540^\circ\text{C}$ ;  $S_t = 8$  Hrs;  $A_T = 180^\circ\text{C}$ ;  $A_t = 14$  Hrs.

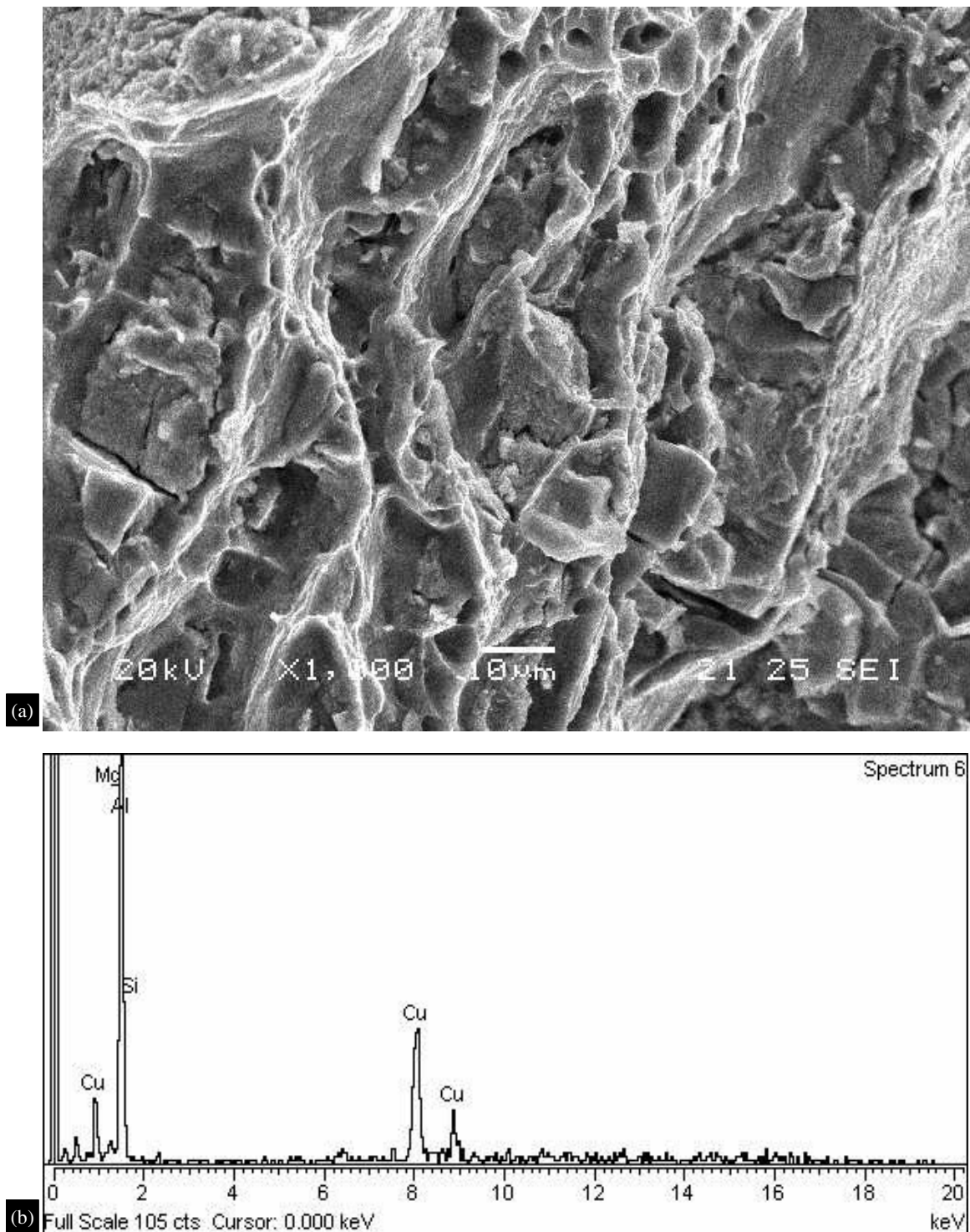
Fracture in engineering alloys can occur via transgranular (across grains) or intergranular (along grain boundaries) fracture paths. However, regardless of the fracture path, there are 4 principal fracture modes (dimple rupture, cleavage, fatigue and decohesive rupture). Each of these modes has a

characteristic fracture surface and a mechanism or mechanisms by which the fracture propagates. To analyze the fracture surface of the specimens to assess the material quality, fractographs of tested specimens were performed using scanning electron microscope (SEM). Energy dispersive X-ray analysis was carried out at different locations of the fractured specimens. EDAX spectrums show chemical composition of inclusions and matrix components. Figure 4 shows the optical microstructures at different heat-treatment processes. SEM of tensile specimens at heat-treatment process ( $S_T = 500^\circ\text{C}$ ;  $S_t = 8$  Hrs;  $A_T = 200^\circ\text{C}$ ;  $A_t = 12$  Hrs) in Figure 5(a) shows combined intergranular decohesion and trans granular cleavage like fracture, while Figure 5(b) for the heat-treatment process ( $S_T = 540^\circ\text{C}$ ;  $S_t = 8$  Hrs;  $A_T = 180^\circ\text{C}$ ;  $A_t = 14$  Hrs) shows transition from intergranular mode to dimple mode of failure, and blight inclusion of SiC (Stringer type) by cleavage. SEM and EDAX spectrum of CT specimens in Figure 6 at  $S_T = 500^\circ\text{C}$ ;  $S_t = 8$  Hrs;  $A_T = 200^\circ\text{C}$ ;  $A_t = 12$  Hrs depicting the failure along intergranular facets (see different sizes of SiC particles).



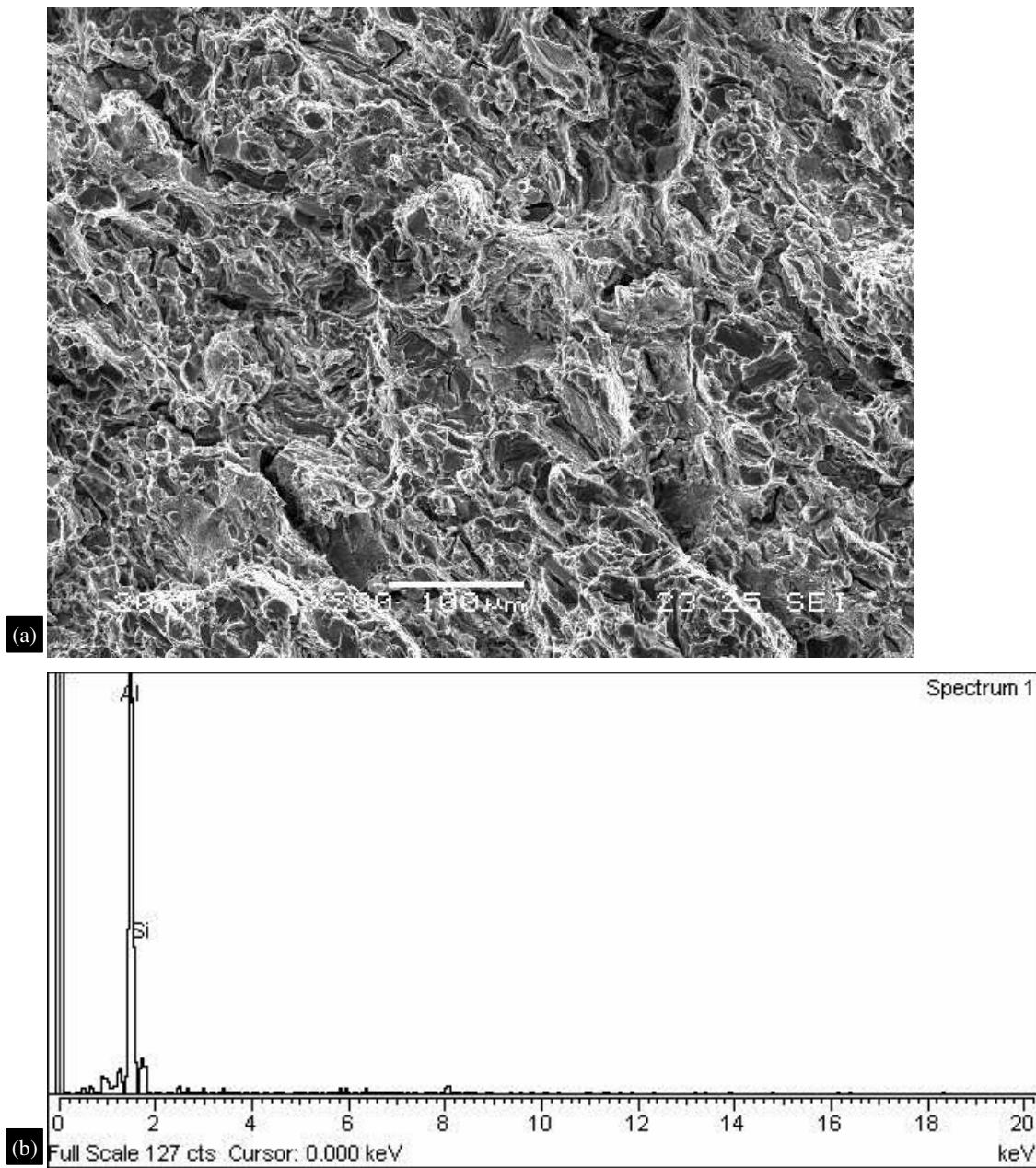
**Figure 5.** SEM of tensile specimens at different heat-treatment processes.



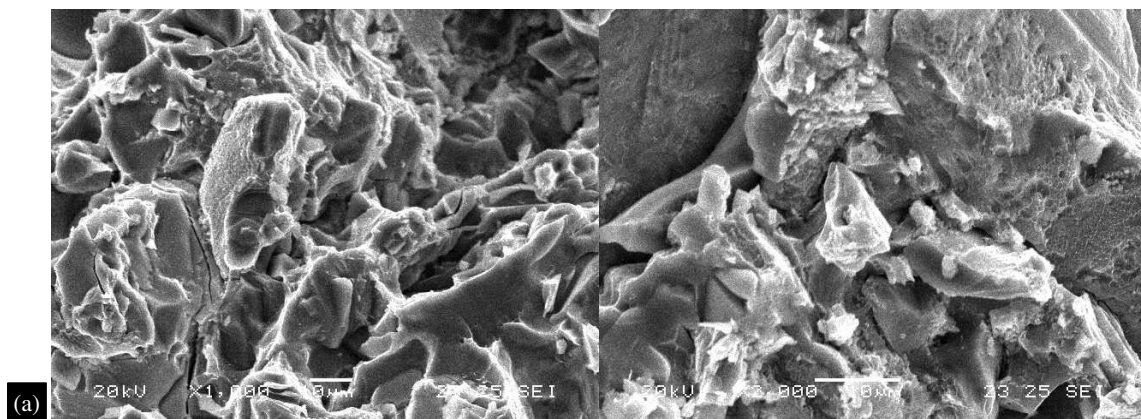


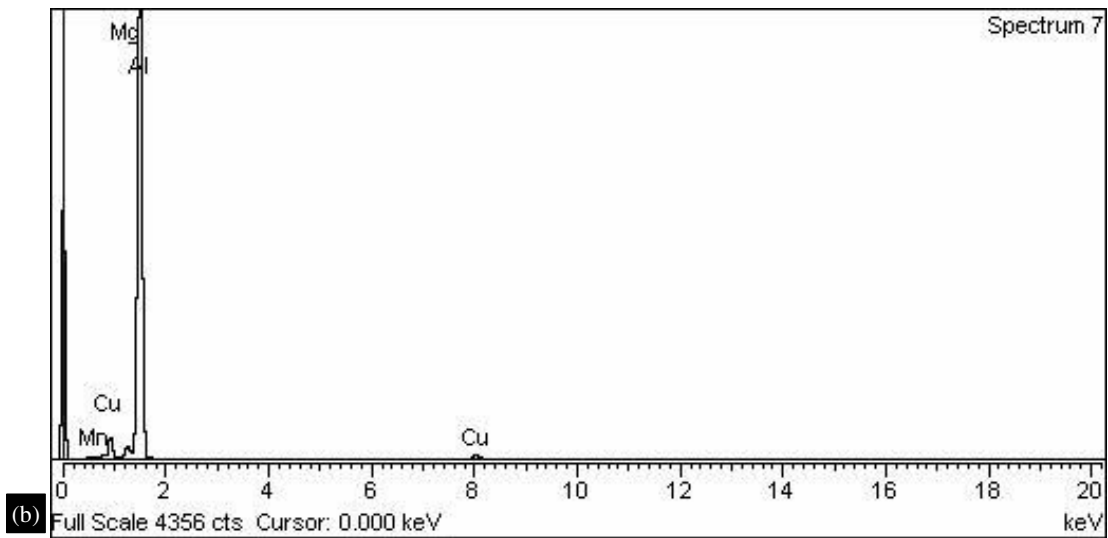
**Figure 6.** SEM and EDAX spectrum of CT specimen at heat-treatment-process:  $S_T = 500^\circ\text{C}$ ;  $S_t = 8$  Hrs;  $A_T = 200^\circ\text{C}$ ;  $A_t = 12$  Hrs.

SEM and EDAX spectrum of CT specimen for the heat-treatment-process ( $S_T = 540^\circ\text{C}$ ;  $S_t = 8$  Hrs;  $A_T = 180^\circ\text{C}$ ;  $A_t = 14$  Hrs) in Figure 7 shows large number of dimples along with intergranular voids (mixed mode of failure). Figure 8 shows the scanning electron micrographs and EDAX spectrum of CT specimens for the hot-pressing parameters: Temperature =  $450^\circ\text{C}$ ; % of compression = 15, whereas Figure 9 shows for the hot-pressing parameters: Temperature =  $500^\circ\text{C}$ ; % of compression = 50.

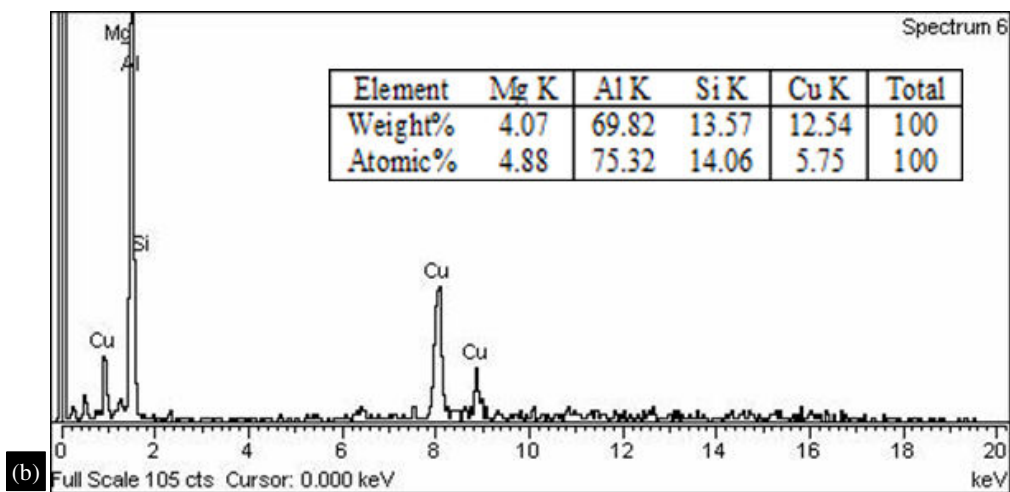
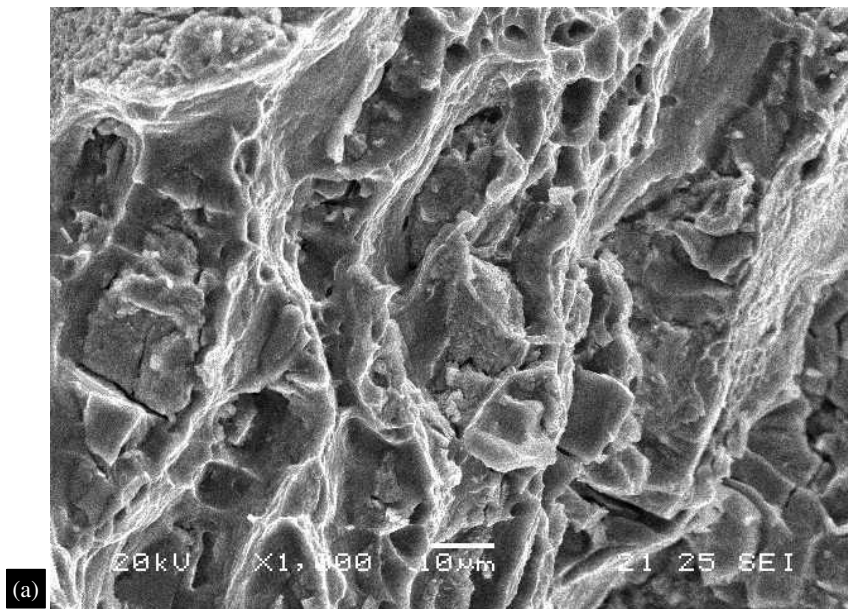


**Figure 7.** SEM and EDAX spectrum of CT specimen for the heat-treatment-process ( $S_T = 540^\circ\text{C}$ ;  $S_t = 8$  Hrs;  $A_T = 180^\circ\text{C}$ ;  $A_t = 14$  Hrs).





**Figure 8.** SEM and EDAX spectrum of CT specimens (Hot pressing parameters:



**Figure 9.** Scanning electron micrographs and EDAX spectrum of CT specimens. (Hot pressing parameters: Temperature = 500°C; % of compression = 50).

## CONCLUSIONS

This paper deals with the fracture behavior of 2124Al-10vol% SiCp composite tensile specimens under different heat treatment processes. Taguchi's  $L_9$  OA (orthogonal array) is selected for the 4 process variables with 3 levels of each variable. The 4 process variables are solutionizing temperature ( $S_T$ ), solutionizing time ( $S_t$ ), aging temperature ( $A_T$ ) and aging time ( $A_t$ ), the effect of which is evaluated on the notched and un-notched tensile strength of the composite. Fracture toughness ( $K_{Ic}$ ) of composites is evaluated by conducting tests on Compact tension (CT) specimens. Scanning electron microscope (SEM) is used to generate factographs on tested tensile and CT specimens. Energy dispersive X-ray analysis was carried out at different locations of the fractured specimens. The chemical composition of the inclusions and matrix components was characterized in EDAX spectrums.

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