# Use of salivary liver function markers for the prognosis of deaddiction in alcohol dependent non-smokers and smokers: A before and after study

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#### Abstract

Alcohol deaddiction improves the liver function markers in blood as well as saliva in both smokers and nonsmokers. The study was conducted to observe the changes in liver function markers after deaddiction in both blood as well as saliva. The study included 83 alcohol dependent subjects, among whom 45 were smokers and 38 were non-smokers. They were followed up after deaddiction for 19-21 days. The liver function markers GGT-  $\gamma$ glutamyl transferase, AST- Aspartate transaminase, ALT- Alanine transaminase, TP- Total protein, and Alb-Albumin were measured spectrophotometrically before and after deaddiction in saliva as well as blood. Paired't' test was done to analyse the significance of their changes in before and after deaddiction. There was no difference in NR (normalization rate) of GGT, AST and ALT between alcohol dependent smokers and non-smokers in saliva as well as blood. NR of TP and Alb was better in non-smokers compared to smokers. GGT (p<.001), AST (p<.001) and ALT (p<.001) have decreased and SFR has increased significantly in all groups after withdrawal. Elevation after withdrawal of TP and Alb were not significant. Elevation of Alb after withdrawal was not significant. In both alcohol dependent smokers as well as non-smokers, salivary GGT (p<.001), AST (p<.001) and ALT (p<.001) levels can be used instead of blood levels to monitor the prognosis of deaddiction, but TP and Alb cannot be used for the same.

Keywords: Aminotransferases, Alcohol withdrawal, Alcohol dependence, y-glutamyl transferase.

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#### Introduction

Alcohol is a psychoactive substance which may cause dependence. The World Health Organization (WHO) recently stated that harmful use of alcohol ranks among the top five risk factors for disease, disability and death throughout the world. Globally, alcohol abuse causes approximately 3.3 million deaths every year or 5.9% of all deaths, and 5.1% of the global burden of disease is attributable to alcohol consumption.<sup>1</sup>

Alcohol dependents are addicted to smoking which may increase the pathogenic effects on the liver.<sup>2</sup> The smoking and alcohol dependence has a common genetic vulnerability.<sup>3</sup>

Alcohol biomarkers play a significant role in the early diagnosis of alcohol abuse, alcoholrelated organ damages, and assessment of alcoholism therapy. The liver function markers assays are used in the detection of excessive alcohol consumption, acute ethanol intoxication and in monitoring of alcohol consumption. Alcohol biomarkers can demonstrate the effects of alcohol on the body, classified as indirect markers, or as direct markers. The traditional alcohol biomarkers are alanine aminotransferase (ALT), aspartate aminotransferase (AST), and Gamma glutamyl transferase (GGT). The activities of GGT, AST and ALT are elevated in chronic  $alcoholics^4$  and indicate generalized hepatic damage.<sup>5</sup>

Saliva like blood contains wide range of enzymes and proteins as biological markers. Blood collection is intrusive however salivary collection is non-invasive and economical.<sup>6</sup> The constituents of the blood enter the saliva by mechanism such as active transport, passive diffusion, leakage or ultrafiltration.<sup>7</sup> Saliva can be used as a substitute for blood in the diagnosis and prognosis of the disease.<sup>8</sup>

Our study hypothesis is that alcohol deaddiction improves the liver function markers in blood as well as saliva in both smokers and non-smokers. The studies to prove the effect of smoking on the alcohol induced changes after deaddiction in liver function enzymes in both blood as well as saliva are rare. Hence the study was conducted to observe the changes in liver function markers after deaddiction in both blood as well as saliva. We also intend to find if there is any difference in normalization of the liver function markers between alcohol dependent smokers and non-smokers.

#### Materials and Methods

The study is a prospective cohort study, conducted at Vailankanni deaddiction centre, Father Muller Medical College, Mangalore, for a period of six months from Jan 2016 to July 2016. The patients admitted for deaddiction were diagnosed as alcohol dependence by a treating psychiatrist based on The ICD-10 (the 10<sup>th</sup> revision of the international statistical classification of diseases) classification of mental and behavioural disorders: diagnostic criteria for research WHO.9 All the subjects were followed up for 21 days after deaddiction for the outcome of normalization in liver function markers. Of the 150 alcohol dependent participants recruited for the study, 83 subjects were found eligible and agreed to participate voluntarily, by providing information about the duration and amount of alcohol consumption, smoking and family history. Based on the history of smoking we categorised alcohol dependents into alcohol dependent smokers and non-smokers. Among them 45 subjects were alcohol dependent smokers with age 36±10 years (mean±SD) ranged from 21-60 years. And 38 subjects were alcohol dependent non-smokers of age 39±10 years (mean±SD) ranged from 23-65 years. The recruited participants with the following potential confounders as in systemic illness such as cancer, thyroid disorder, alcohol liver disease, inflammatory disease, psychiatric illness, abstinence from alcohol prior to admission, and substance other than alcohol and nicotine dependence were excluded from the study. The written informed was consent was taken from the study subjects. This study was approved by Father Muller Institutional Ethics committee.

For collection of unstimulated whole saliva, subjects were refrained from taking food for 1 hour before collection and advised to clean the mouth by rinsing with water to remove any food debris. The saliva was collected between 9AM-11AM into a sterile containers avoiding forcible spitting as specified by the method of Navzesh.<sup>10</sup> The time and amount of saliva collected is recorded for the calculation of salivary flow rate (SFR). The saliva samples were centrifuged at 3000 rpm for 15 minutes and the supernatant was collected. Blood was collected after the saliva collection in EDTA or plain vaccutainers taking aseptic precautions, serum is separated by centrifugation. The saliva and serum samples collected were stored in  $-80^{\circ}$  C until the assay done.

Biochemical assays for saliva and serum GGT, AST, ALT, TP Alb, and Glb (globulin) were performed in all subjects. GGT activity was assayed by carboxy substrate method.<sup>11</sup> Amino transferases activity both AST<sup>12</sup> and ALT<sup>13</sup> were assayed by modified IFCC method. Albumin was assayed by Bromocresol green dye binding method.<sup>14</sup> The salivary TP was assayed by Lowry's method<sup>15</sup> and serum TP was assayed by Biuret's reagent method.<sup>16</sup> Glb was calculated by TP-Alb. Paired 't' test was employed to assess the changes in salivary and blood liver function markers before and after withdrawal. Paired correlation was done to assess the strength of association between before and after withdrawal levels of liver function markers. SPSS version 24 software was used for the analysis. p value <.05 was considered statistically significant.

#### Results

At the time of study the subjects recruited was 83(45 alcohol dependent smokers and 38 alcohol dependent non-smokers) there was no loss to follow up. The descriptive data is shown in Table 1. Results of paired't' test of salivary and blood liver function markers before and after withdrawal in alcohol dependent smokers and non-smokers are shown in Table 2 and 3 respectively.

The correlation of paired samples of salivary and blood liver function markers in alcohol dependents, smokers and non-smokers are shown in Table 4 and 5 respectively.

Table 1: Demographic variables of the study population

Demographic variables	Alcohol dependents (both smokers + non-smokers) (n-83)				
Age in years{Mean±SD(range)}	37.5±10(21-65)				
Duration of alcohol consumption in years Median (IQR)	15(8)				
Amount of alcohol consumption in ml	<180ml-n=18, 180-360ml-n=37,				
per day	360-540ml-n=20, >540ml-n=8.				
Family history of Alcohol dependence	Present – 36(43%), Absent- 47(56%)				
History of smoking present	n-45				
History of smoking absent	n-38				
Daily consumption in years	6.61				
Morning consumption in years	3.75				

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<b>G P</b>	COT						CIII
Salivary	GGT	AST (IU/L)	ALT	TP (g/dl)	Alb (g/dl)	SFR (ml/min)	Glb
Liver	(IU/L)	Mean	(IU/L)	Mean	Mean	Mean	(g/dl)
function	Mean	difference	Mean	difference	difference	difference	Mean
markers	difference	±SE	difference	±SE	±SE	±SE	difference
	±SE	(95%CI)	±SE	(95%CI)	(95%CI)	(95%CI)	±SE
	(95%CI)		(95%CI)				(95%CI)
Alcohol	4.32±.37	6.07±.52	3.51±.4	$004 \pm .009$	$01 \pm .005$	.05±.01	.001±.001
dependents	(3.5-5)	(5.04-7.1)	(2.7-4.3)	(0201)	(0200)	(0803)	(0103)
	*p-<.001	*p-<.001	*p-<.001	p59	p05	*p-<.001	p511
	NR-48.3%	NR-48.8%%	NR-43.4%	NR-(- 3.38%)	NR-(-28.1%)	NR- (-18.9%)	NR-(-6.3%)
Alcohol	$4.08 \pm .5$	6.5±.79	3.19±.54	.001±.01	$001 \pm .003$	$05 \pm .01$	.003±.01
dependent	(3.06-5.1)	(4.9-8.1)	(2.1-4.2)	(026028)	(008005)	(0802)	(0203)
smokers	*p-<.001	*p-<.001	*p-<.001	p93	p64	*p001	p844
	NR-48.1%	NR-49.2%	NR-41%	NR- 0.72%	NR-(-3.8%)	NR-(-18.3%)	NR-(-2.3%)
Alcohol	4.6±.55	5.5±.63	3.8±.61	$01 \pm .01$	$02\pm.01$	$06 \pm .02$	.01±.01
dependent	(3.4-5.7)	(4.2-6.8)	(2.6-5.1)	(0301)	(040006)	(10-(02))	(0204)
non-	*p-<.001	*p-<.001	*p-<.001	p31	p05	*p004	p466
smokers	NR-48.4%	NR-48.08%	NR-45.8%	NR-(-8.84%)	NR-(-54.3%)	NR-(-19.5%)	NR- (-12%)

Table 2: Paired't' test of salivary liver function markers before and after withdrawal in alcohol dependent smokers and non-smokers

Legend to Table 2: GGT-  $\gamma$  glutamyl transferase, AST- Aspartate transaminase, ALT- Alanine transaminase, TP- Total protein, Alb-Albumin, SFR- Salivary flow rate, SE- Standard error, CI-Confidence interval, NR- Normalization rate, p<.05 is statistically significant. NR = (Levels before withdrawal-Level after withdrawal) X100/Level before withdrawal. There was no difference in NR of GGT, AST ALT and SFR between the groups. NR of TP and Alb was better in non-smokers compared to smokers. GGT, AST and ALT have decreased and SFR has increased significantly in all groups after withdrawal. Elevation after withdrawal of TP, Alb and Glb were not significant. p<.05 – significant.

Table 3: Paired't' test of Blood liver function markers before and after withdrawal in alcohol dependent smokers and non-smokers

Blood Liver function markers	GGT (IU/L) Mean difference ±SE(95%CI)	AST (IU/L) Mean difference ±SE(95%CI)	ALT (IU/L) Mean difference ±SE(95%CI)	TP (g/dl) Mean difference ±SE(95%CI)	Alb (g/dl) Mean difference ±SE(95%CI)	Glb (g/dl) Mean difference ±SE(95%CI)
Alcohol	214.8±33.3	70.3±7.5	43.6±5	2±.06	$04 \pm .05$	2±.05
dependents	(148-281)	(55.4-85)	(33.6-53.7)	(34-(06))	(1505)	(3 – (05))
	*p-<.001	*p-<.001	*p-<.001	*p004	p352	*p005
	NR- 66.7%	NR- 61.83%	NR-55.7%	NR- (-2.7%)	NR-(-1.1%)	NR- (-5.3%)
Alcohol	264±54	70.9±11.4	41.3±6.8	$14\pm.09$	$02\pm.07$	012±.06
dependent	(153-374)	(47.9-94)	(27-55)	(3304)	(171)	(301)
smokers	*p-<.001	*p-<.001	*p-<.001	p14	p69	p078
	NR- 70.5%	NR- 60.1%	NR- 52.1%	NR- (-1.9%)	NR- (-0.7%)	NR- (-3.9%)
Alcohol	156±31.9	69.6±9.4	46.3±7.6	27±.09	07±.07	2±.09
dependent	(91.8-221)	(50.4-88.7)	(30.9-61.8)	(47(07))	(2207)	(4 –(-0.02))
non-	*p-<.001	*p-<.001	*p-<.001	*p007	p345	*p03
smokers	NR- 59.9%	NR-63%	NR- 60%	NR- (-3.7%)	NR-(-1.6%)	NR- (-6.9%)

Legend to table No.3: GGT-  $\gamma$  glutamyl trasnferase, AST- Aspartate transaminase, ALT- Alanine transaminase, TP- Total protein, Alb-Albumin, SFR- Salivary flow rate, SE- Standard error, CI-Confidence interval, NR- Normalization rate, p<.05 is statistically significant. NR = (Levels before withdrawal-Level after withdrawal) X100/Level before withdrawal. There was no difference in NR of GGT, AST and ALT between the groups. NR of TP and Alb was better in non smokers compared to smokers. GGT, AST and ALT have decreased significantly in all groups after withdrawal. TP elevated significantly after withdrawal in non-smokers. Elevation of Alb after withdrawal was not significant. Glb was significantly decreased in smokers after withdrawal. p<.05 – significant.

 Table 4: Paired samples Correlation of salivary liver function markers in alcohol dependents, smokers and non-smokers

	Alcohol dependent			Alcohol dependent smokers			Alcohol dependent non-smoker		
	Mean± SE BW	Mean± SE AW	r value P value	Mean± SE BW	Mean± SE AW	r value P value	Mean± SE BW	Mean± SE AW	r value P value
GGT	8.9±	4.65±	0.792	8.5±	4.4±	0.77	9.56±	4.96±	0.81
(IU/L)	0.57	0.33	*p<.001	0.78	0.45	*p<.001	0.87	0.49	*p<.001
AST	12.45±	6.37±	0.833	13.24±	6.7±	0.8 *p<.001	11.5±	5.97±	0.88

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	0.9	0.59	*n< 001	1.25	0.72		13	0.99	*n< 001
ALT	8.18+	4.6+	0.843	7.84+	4.64+	0.78	8.6+	4.7+	0.9
(IU/L)	0.65	0.34	*p<.001	0.82	0.46	*p<.001	1.04	0.52	*p<.001
ТР	0.14±	0.15±	0.323	0.153±	0.152±	0.32	0.14±	0.15±	0.32
(g/dl)	0.008	0.006	*p=.003	0.013	0.009.	*p<.029	0.01	0.01	*p=.048
Alb	$0.04\pm$	$0.05 \pm$	0.376	0.039±	0.041±	0.74	0.04±	0.06±	0.26 - 11
(g/dl)	0.003	0.006	*p<.001	0.005	0.004	*p<.001	0.005	0.01	0.20 p=.11
Glb	0.1±	0.09±	0.327	0.113±	0.11±	0.376	0.09±	$0.08\pm$	0.27
(g/dl)	0.009	0.008	*p=.003	0.014	0.009	*p=.011	0.01	0.014	p=.097
SFR	0.30±	0.36±	0.758	0.29±	0.35±	0.83	0.32±	0.38±	0.68
(ml/min)	0.018	0.014	*p<.001	0.026	0.019	*p<.001	0.03	0.02	*p<.001

Legend to Table 4: GGT-  $\gamma$  glutamyl trasnferase, AST- Aspartate transaminase, ALT- Alanine transaminase, TP- Total protein, Alb-Albumin, SFR- Salivary flow rate, SE- Standard error. BW-Before withdrawal, AW- After withdrawal, r-correlation coefficient. All the parameters show significant correlation between before and after withdrawal levels except TP, Alb and Glb in alcohol dependent non-smokers. p<.05 – significant.

 Table 5: Paired samples Correlation of blood liver function markers in alcohol dependents, smokers and non-smokers

	Alc	ohol deper	Ident	Alcohol dependent smokers			Alcohol dependent non-smoker		
	Mean± SE BW	Mean± SE AW	r value p value	Mean± SE BW	Mean± SE AW	r value p value	Mean± SE BW	Mean± SE AW	r value p value
GGT (III/L)	322±	107±	0.772	374.4±	110.3±	0.697	261.4±	104.9±	0.95
001 (IC/L)	41	11	*p<.001	64	64 14.8	*p<.001	47.6	16.8	*p<.001
AST	113±	43.4±	0.710	116.6±	45.6±	0.686	110.5±	40.9±	0.761
(IU/L)	9.4	3.2	*p<.001	14.4	5.1	*p<.001	12.1	4	*p<.001
ALT	78.4±	34.7±	0.766	79.4±	38.1±	0.712	77.3±	30.1±	0.837
(IU/L)	6.9	2.9	*p<.001	9	4	p<.001	10.8	4.2	p<.001
ТР	7.4±	7.6±	0.274	7.4±	7.5±	0.179	7.4±	7.7±	0.376
(g/dl)	0.05	0.05	*p=.012	0.08	0.07	p=.24	0.09	0.08	*p=.02
Alb	4.42±	$4.47\pm$	0.105	4.4±	4.5±	0.05	4.4±	4.5±	0.172
(g/dl)	0.04	0.03	p=.343	0.05	0.05	p=.74	0.06	0.05	p=.303
Glb	2.98±	3.1±	0.475	2.96±	3.1±	0.496	3±	3.2±	0.46
(g/dl)	0.05	0.05	*p<.001	0.07	0.06	*p=.001	0.01	0.01	*p=.004

Legend to Table 5: GGT-  $\gamma$  glutamyl trasnferase, AST- Aspartate transaminase, ALT- Alanine transaminase, TP- Total protein, Alb-Albumin, SE- Standard error. BW- Before withdrawal, AW-After withdrawal, r-correlation coefficient. GGT, AST, ALT, and Glb show significant correlation between before and after withdrawal levels. TP was significant in alcohol dependent smokers in both before and after withdrawal. Alb was not significant in any groups. p<.05 – significant.

#### Discussion

The significant decrease observed in our study in the levels of GGT, AST and ALT were not only in alcohol dependent non-smokers but also in alcohol dependent smokers. This finding was seen in blood as well as saliva. As observed in our study, several studies have corroborated our finding that even though smoking enhances the effect of alcohol it does not cause direct liver injury.<sup>17</sup> Alcohol leads to inflammation with increase in the inflammatory liver enzymes GGT, AST and ALT levels as observed in our study.<sup>18</sup> However after withdrawal we observed significant normalization in the liver enzymes, this could be due to the significant recovery from the alcohol induced hepatic inflammatory activity.

The normalization rates of liver enzymes in this study after deaddiction were very similar in blood as well as saliva which is concordant with other studies.<sup>19</sup> Hence salivary liver function enzymes can be used to monitor the prognosis of the deaddiction treatment in both alcohol dependent smokers as well as non-smokers. However large population based studies are needed to validate this.

The recovery from the liver inflammation usually leads to improvement in synthetic functions of the hepatocytes.<sup>20</sup> This is responsible for the elevated levels of the TP and Alb in saliva as well as blood as observed in our study but it was not significant. In our study the after withdrawal levels of total protein and albumin were measured varying from 19-21 days of after deaddiction, which is almost equal to the half life of albumin.<sup>21</sup> This clearly indicates hepatocytes require longer time to regain their synthetic activity. The expected elevation in TP levels after withdrawal was not as anticipated because of the concomitant decrease in the globulin levels, which could be due to recovery from the alcohol induced hepato inflammatory activity.

The decrease observed in Glb after withdrawal was significant only in blood of alcohol dependent non-smokers. The additive enhancing inflammatory effect of smoking is responsible for the persistent elevation of the globulins after withdrawal in alcohol dependent smokers (Table 2 and 3).

Total protein and albumin levels after withdrawal cannot be predicted using values of before withdrawal levels in blood as well as saliva in both alcohol dependent smokers and non-smokers. The increase in total protein and albumin observed after withdrawal is not uniform with respect to their before withdrawal levels (Table 4 and 5).

Further studies are needed to prove this hypothesis that hepato inflammatory activity recovers earlier than the regain in the hepatic synthetic activity after deaddiction. Serum and salivary liver enzymes should be correlated with hsCRP and other inflammatory markers.

#### Conclusion

In both alcohol dependent smokers as well as non-smokers, salivary GGT, AST and ALT levels can be used instead of blood levels to monitor the prognosis of deaddiction, but TP and Alb cannot be used for the same.

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### Compliance with ethical standards

**Conflict of Interest:** JB Honnamurthy, AR Shivashankara, SS Avinash, P JohnMathai, and M Malathi declare that they have no conflict of interest.

## Human and Animal Rights Statement: All procedures performed in studies involving

human participants were in accordance with the ethical standards of the institutional research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

**Ethical approval:** The study was approved by Father Muller Institutional ethics committee. (FMMC/FMIEC/2039/2014).

**Informed consent:** All authors declare that written informed consent was obtained from all individual participants included in the study.

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