Original article

Role of maternal serum inhibin A and insulin – like growth factor 1 levels as predictors of fetal outcome in high risk pregnancies.

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Abstract

Background: Timely prediction of adverse pregnancy and perinatal outcome has been the endeavor of clinicians since a long time. Identification and validation of a serum based biomarker is the need of the hour and many tentative molecules have been studied with variable results. Two such molecules are inhibins and insulin like growth factors. This study was planned to assess the utility of inhibin A and IGF-1 as predictors of adverse pregnancy and fetal outcomes.

Methods: The study population comprised of all pregnant women presenting to the ante natal clinic in SSK hospital between gestation periods of 24-28 weeks fulfilling the inclusion criteria. The criteria adopted for inclusion in the study included primi-gravida with singleton pregnancy or multi-gravida with high risk factors in the previous pregnancy such as gestational hypertension, preeclampsia, eclampsia, placental abruption, intrauterine growth retardation, intrauterine fetal demise, preterm delivery and early pregnancy loss.

Results: The mean levels of inhibin A were significantly higher in women with overall adverse pregnancy outcome such as preeclampsia, gestational hypertension, and preterm delivery as compared to the normal counterparts. No significant difference in the level of IGF-1 was observed with various adverse outcomes. Inhibin A at the cutoff of > 443pg/ml was found to be reliable marker of preeclampsia and preterm delivery.

Conclusion: We may hypothesize from our findings that second trimester inhibin A levels may be further evaluated through large prospective studies as an reliable biomarker of adverse pregnancy outcome. *Keywords: Preeclampsia, IGF-1, Inhibin A, SGA*

Introduction

Preeclampsia is a common hypertensive disorder affecting 7 -10 % of pregnancies with profound implications for both maternal and fetal well being (1). The in depth understanding of the various molecular pathways that predispose certain women to develop this condition is still vague. According to the present understanding, preeclampsia develops as a two stage process with poor placentation in the first half of pregnancy followed by maternal response in the second half of pregnancy(2).

Although medical science has made great progress in the recent past, accurate prediction of adverse pregnancy outcomes such as preeclampsia, preterm deliveries, intrauterine growth retardation, still births etc still elude clinicians. They contribute greatly to maternal and fetal morbidity and mortality. Identification and validation of a serum based biomarker is the need of the hour and many tentative molecules have been studied with variable results. Two such molecules are inhibins and insulin like growth factors.

Inhibins are dimeric glycoproteins which are members of the transforming growth factor β superfamily(3). Inhibin A is produced by the placenta and has been shown by some researchers to be raised as early as 10-15 weeks of pregnancy in women who subsequently develop preeclampsia (4). Insulin like growth factor-1 (IGF-1) is a mitogenic polypeptide that is thought to play a critical role in abnormal placentation- a hallmark of preeclampsia (5).

This study was planned to assess the utility of inhibin A and IGF-1 as predictors of adverse pregnancy and fetal outcomes.

Method

The study was conducted by the collaboration between the departments of biochemistry and obstetrics and gynecology, Lady Hardinge Medical College and associated SSK hospital, New Delhi. The study was commenced after procuring due clearance from the institutional ethical committee. Bilingual written informed consent was taken from all eligible subjects with detailed explanation of methodology, purpose and implicit risks if any. A total of 60 subject were enrolled in the study.

The study population comprised of all pregnant women presenting to the ante natal clinic in SSK hospital between gestation periods of 24-28 weeks fulfilling the inclusion criteria. The criteria adopted for inclusion in the study included primi-gravida with singleton pregnancy or multi-gravida with high risk factors in the previous pregnancy such as gestational hypertension, preeclampsia, eclampsia, placental abruption, intrauterine growth retardation, intrauterine fetal demise, preterm delivery and early pregnancy loss. Subjects with chronic hypertension, renal failure, Diabetes Mellitus, thyroid disorders, multiple pregnancies, epilepsy and fetal congenital malformations were excluded from the study.

After detailed clinical examination and routine investigations, the subjects were called for regular follow up visits. Blood was obtained by venepuncture and the serum stored at -20 degree Celsius for special investigations. Serum inhibin A and IGF-1 levels were assayed by commercially available ELISA kits supplied by DRG international, Germany. The inter-assay and intraassay precision of Inhibin A was 7.3% and 3.6 % respectively. While the same values for IGF-1 were 7.2% and 4.2% respectively.

The maternal outcome was assessed by the development of gestational hypertension, placental abruption, preeclampsia and eclampsia while the

fetal outcome was gauged by the mode of delivery, period of gestation at the time of delivery, birth weight, APGAR score at 5 min, need for neonatal resuscitation or admission to neonatal ICU, small for gestational age babies and intrauterine fetal death or early neonatal death.

Statistical analysis

All data analysis was done using SPSS package version 19 (SPSS Inc., Chicago, IL, USA). Data were presented as mean \pm standard deviation (standard error of means) or number of subjects and percentage. Chi- square test or student's t test was used for comparison of means between parametric data. The Mann Whitney test was used to compare the means in case of non parametric data. The sensitivity and specificity of various markers ass determinants of risk were determined. P value <0.05 was considered significant.

Results

The demographic profile of the study population is tabulated in table no 1. The mean age of the subjects was 26.38 ±4.82 yrs with the maximum number (29/60) participants were in the 21-25 yrs age group. The mean BMI of the study population at the time of enrollment was $23.23 \pm 4.00 \text{ kg/m}^2$. The mean gravid status of the subjects was 2.5 ±1.3. Approximately 20% of the population were primigravida, 45% were of gravid 2, 23.3% were of gravid 3 and 11.7% were of gravid \geq 4. The mean period of gestation at the time of enrollment was 23.5 ± 1.35 weeks.

Parameter	Mean ± SD
Age (yrs)	26.38 ±4.82
BMI (Kg/m ²)	23.23 ± 4.00
Gravida	2.5 ±1.3
Period of gestation (weeks)	23.5 ± 1.35

Table 2: The distribution of risk factors is depicted in table

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Risk factor	Number	Percentage
Primigravida	12	20%
Previous history of		
Preeclampsia	16	26.7%
Eclampsia	02	3.3%
Gestational	09	15%
Hypertension		
Placenta Abruption	02	3.3%
Intrauterine Death	03	5%
Intrauterine Growth	09	15%
restriction		
Preterm Delivery	11	18.3%
Early Pregnancy Loss	22	36.7%

Primigravida comprised of 20 % of the study population whereas preeclampsia was present in 27 % of the previous pregnancies of the multi-parous women.

The maternal outcome pattern was also studied. Normal maternal outcome was observed in 52 subjects whereas adverse maternal outcome in the form of gestational hypertension, preeclampsia, eclampsia, placental abruption was observed in 8 subjects. Vaginal delivery occurred in 51 pregnancies whereas LSCS was conducted in 9 pregnancies in the study population.

The fetal outcome is illustrated in table no 3. The mean birth weight was 2.58 ± 0.55 kg while the mean gestational age was 37.50 ± 1.97 weeks. The other parameters which were assessed include SGA, Apgar score, neonatal resuscitation and admission to neonatal ICU.

Fetal outcome	Number	Percentage (%)				
Birth Weight (Kg)						
< 1.5 Kg	1	1.7				
1.5 – 1.9 Kg	5	8.3				
2.0 – 2.4 Kg	15	25				
> 2.5 Kg	39	65				
Gestational Age at I	Delivery					
< 34	3	5				
34 - 36	13	21.7				
> 36	44	73.3				
SGA						
< 10 th centile	8	13.3				
10 -90 th centile	52	86.7				
>90 th centile	0					
APGAR score at 5 n	ninutes					
<7	5	8.3				
>7	55	91.7				
Neonatal	1	1.7				
Resuscitation						
Admission to	2	3.3				
neonatal ICU						

The serum levels of inhibin A and IGF-1 in pregnancies with normal and adverse outcomes are compared in tables 4a and 4b respectively.

The mean levels of inhibin A were significantly higher in women with overall adverse pregnancy outcome such as preeclampsia, gestational hypertension, and preterm delivery as compared to the normal counterparts.

	Advers	e	Normal outcome		_	
	outcom	ne				Р
	Mean		Mean inhib	Mean inhibin A		
	inhibin	А				
	Pg/m	Mo		Pg/m	MoM	
	1	Μ		1		
Overall	443.	1.9	Overall	189.	$0.85\pm$	< 0.0
adverse	94±	$9 \pm$	normal	42±	0.11	01
outcome	57.7	0.2	outcome	24.8		
(N=24)	4	6	(N=36)			
Preecla	632.	2.8	Normote	235.	$1.06 \pm$	0.00
mpsia	7 ±	$4 \pm$	nsive	74±	0.12	6
(N=4)	143.	0.6	(N=52)	27.0		
	48	4		7		
Gestatio	395.	1.7	Normote	235.	$1.06 \pm$	0.06
nal	$98 \pm$	$8 \pm$	nsive	74±	0.12	5
hyperten	72.3	0.3	(N=52)	27.0		
sion		3		7		
(N=4)						
Preterm	440.	1.9	Term	211.	0.95	0.00
Delivery	$49 \pm$	$8 \pm$	delivery	95	±0.11	3
(N=16)	70.1	0.3	(N=44)	±24.		
	2	1		21		
SGA	287.	1.2	AGA	270.	1.22	0.94
(N=8)	$81 \pm$	9 ±	(N=52)	60 ±	±0.14	8
	85.9	0.3		30.6		
	9	8		2		
Low	203.	0.9	Normal	279.	1.26±	0.74
APGAR	$23 \pm$	$1 \pm$	APGAR	23	0.14	8
score	58.7	0.2	(N=55)	±30.		
(N=5)	3	6		75		

Р
val
ue
0.0
63
0.1
92
0.7
38
0.0
90
0.2
07
0.4
22

The mean IGF-1 levels in the cases that developed adverse outcome were not significantly lower than those with normal pregnancy outcome. No

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significant difference in the level of IGF-1 was observed with various adverse outcomes.

Table no 5 is the comparative tabulation of inhibin
A and IGF-1 as predictors of adverse pregnancy
outcomes. The cut off for Inhibin A was taken as

443 pg/ml while that for IGF-1 was fixed at 168ng/ml.

outcome	Number of patients With the cut off inhibi	Sensitivity n	Specificity	P value
Normal(n=52)	< 443 43 > 443 9	75	82.7	0.028
Preeclampsia(n=4)	$ \begin{array}{c c} \hline < 443 & 1 \\ \hline \geq 443 & 3 \\ \end{array} $	_		
Normal(n=52)	$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	50	80.4	0.169
Gestational hypertension (n=4)	< 443 2 $\ge 443 2$			
Preterm babies(n=16)	< 443 9 ≥ 443 7	43.8	84.1	0.038
Term babies (n=44)	< 443 7 ≥ 443 37			
SGA(n=8)	$< 443 \qquad 6$ $\ge 443 \qquad 2$	25	76.9	1.000
AGA (n=52)	< 443 40			
Low APGAR score babies (n=5)	$< 443 \qquad 4$ $\ge 443 \qquad 1$	20	76.4	1.000
Normal APGAR score babies (n=55)	$< 44\overline{3}$ 42 ≥ 443 13	_		

Inhibin A at the cutoff of \geq 443pg/ml was found to be reliable marker of preeclampsia and preterm delivery. However it was not significant for gestational hypertension, SGA and low APGAR score.

outcome	Number of patients With the cut off IGF-1		Sensitivity	Specificity	P value
Normal(n=52)	<u><</u> 168	22	75	57.7	0.314
	>168	30			
Preeclampsia(n=4)	<u><</u> 168	3			
	>168	1			
Normal(n=52)	<u><</u> 168	22	50	78.6	1.000
	> 168	30			
Gestational hypertension	<u><</u> 168	2			
(n=4)	> 168	2			
Preterm babies(n=16)	<u><</u> 168	10	62.5	61.4	0.144
	> 168	6			
Term babies (n=44)	<u><</u> 168	17	1		

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	>168	27			
SGA(n=8)	<u><</u> 168	5	62.5	57.7	0.448
	>168	3			
AGA (n=52)	<u><</u> 168	22			
	>168	30			
Low APGAR score	<u><</u> 168	4	80	58.2	0.164
babies (n=5)	> 168	1			
Normal APGAR score	<u><</u> 168	23			
babies (n=55)	>168	32			

Discussion

The present study was carried out to assess the predictive value of second trimester Inhibin A and IGF-1 values as biomarkers for evaluation of risk for adverse outcome pregnancies. Adverse outcome was assessed by the appearance of gestational hypertension, pre eclampsia during pregnancy and preterm delivery, SGA and low APGAR score during parturition.

In the present study, 40% women developed adverse outcome with gestational hypertension and preeclampsia present in 5 and 6.7% of them. Similar figures for incidence of hypertensive disorders of pregnancy were reported by Aquilina et al (6) and Aye et al (7). Approximately 27% of the women later developed preterm delivery while 35% had low birth weight babies. Dane et al (8) and Stepan et al (9) found comparable demography in their study cohorts as well.

Inhibin A levels rises with gestational age during pregnancy. The increase has been attributed to the high prostaglandin and oxytocin levels during pregnancy (10). Studies have shown association with high inhibin A levels with incidence of trisomy 21(11). Furthermore many researchers have now also hinted towards the association of inhibin A >2.0 MoM with adverse pregnancy outcomes (12). Our study also illustrates the same finding.

Inhibin A levels were higher in gestational hypertension as well as in pre eclampsia but statistically significant difference was observed in preeclampsia only. Similar results were echoed in studies conducted by Spencer et al (13) and Florio et al (14). Aye et al found 3.26 MoM in women who developed preeclampsia and 0.99 in women who remained normotensive. Corresponding values form our study were 2.84 MoM and 1.11 MoM respectively(7).

A similar statistically significant difference was observed in case s with pre term delivery as compared to term delivery. Fitzgerald et al reported that higher inhibin A levels increase the risk for intra uterine growth retardation in the fetus(15). Our study also demonstrated higher values in SGA although the difference form AGA was not statistically significant. Daval et al (16)demonstrated that a significant positive correlation can be discerned between second trimester serum inhibin A levels and the development preeclampsia.

Insulin like growth factor-1 (IGF-1) is similar in structure to insulin. IGFs are important determinants of fetal growth and postnatal development (17). IGF-1 is a mitogenic polypeptide that stimulates cellular proliferation and differentiation. Hernandez- Valencia et al showed that lower IGF-1 levels during pregnancy are associated with intra uterine growth retardation (18). IGF system is also thought to play a role in abnormal placentation thereby highlighting its role in pre eclampsia (19).

In the present study, the IGF-1 levels were lower in women who developed preeclampsia as compared to nomotensive controls. However the decrease was not statistically significant. Our findings are concordant with observations reported by Giudice et al (19), Halhali et al (20). Similarly, lower IGF-1 levels were found in gestational hypertension in our study. Shang et al reported significantly lower IGFpregnancies complicated levels in by hypertension (21). Statistically insignificant reduction in IGF-1 levels was observed in pregnancies complicated by adverse fetal or perinatal outcomes. Chia et al also reported significantly reduced IGF-1 levels in IUGR pregnancies leading to SGA babies as compared to AGA babies (22). Lindsay et al concluded from their study on pregnant women with gestational Diabetes Mellitus that fetal IGF-1 levels were inversely associated with maternal diabetes (23).

Comparative assessment of the two markers as indicators of adverse pregnancy outcome revealed that inhibin levels higher than the cutoff of 443pg/ml significantly predicted the occurrence of preeclampsia and preterm delivery. However IGF-1 levels did not show any statistically significant role as an reliable predictor of adverse pregnancy and fetal outcome. We may hypothesize from our findings that second trimester inhibin A levels may be further evaluated through large prospective studies as an reliable biomarker of adverse pregnancy outcome.

References

1. Viler K. Eclampsia and pre-eclampsia: a health problem for 2000 years. In: Critchley H, Maclean A, Poston L,editors. Preeclampsia. London, UK: RCOG press;2003:189-207

2. Redman CW, Sargent IL. Latest advances in understanding preeclampsia. Science 2005;308(5728):1592-4.

3. Muttukrishna S, George L, Fowler PA, Groome NP, Knight PG. measurement of serum concentrations of inhibin A during human pregnancy. Clin Endocrinol 1995;2:391-97.

4. Muttukrishna S, North RA, Morris J, Schellenberg JC, Taylor RS, et al. serum inhibin A abd activin A are elevated prior to onset of preeclampsia. Hum Reprod 2000;15(7):1640-5.

5. Giudice LC, Martina NA, Crystal RA, et al. insulin like growth factor binding protein-1 at the maternal-fetal interface and insulin – like growth factor –I, insulin- like growth factor –II, and insulin-like growth factor binding protein-I in the circulation of women with severe preeclampsia. Am J Obstet Gynecol 1997;176:751-58.

6. Aquilina J, Barnett A, Thompson O, Harrington K. comprehensive analysis of uterine artery flow velovity waveforms for the prediction of preeclampsia. Ultrasound Obstet Gynecol 2000;16: 163-170.

7. Ay E, Kavak ZN, Elter K, Gokaslan H, Pekin T. Screening for preeclampsia by using maternal serum inhibin A, activin A, human

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chorionic gonadotropin, unconjugated estriol, and alpha fetoprotein levels and uterine artery Doppler in the second trimester of pregnancy. Aust N Z J Obstet Gynecol 2005;45(4):283-8.

8. Dane B, Batmaz G, Kilavuz K, Rustemoglu Y, Guler H, Dane C. analysis of the relationship between maternal second trimester AFP, HCG, estriol levels and uterine artery Doppler findings in the prediction of pregnancy complications. Perinatal Journal 2012;20(1):24-29.

9. Stepan H, Unversucht A, Wessel N, Faber R. Predictive value of maternal angiogenic factors in second trimester pregnancies with abnormal uterine perfusion. Hypertension 2007;49(4):818-24.

10. Lockwood CJ. Predicting premature delivery- not easy task. N Engl J Med 2002;364(4):282-4.

11. Aitken DA, Wallace EM, Crosley JA, Swanston IA, van Pareren Y, van Maarle M et al. Dimeric Inhibin A as a marker for Down's syndrome in early pregnancy. N Engl J Med 1996;334(19):1231-6.

12. Dugoff L, Hobbins JC, Malone FD, Vidaver J, Sullivan L, Canick JA et al. FASTER Trial Research Consortium. Quad screen as apredictor of adverse pregnancy outcome. Obstet Gynecol 2005;106(2):260-7.

13. Spencer K, Yu CK, Savvidou M, Papgeorghiou AT, Nocolaides KH. Prediction of preeclampsia by uterine artery Doppler ultrasonography and maternal serum pregnancy-assocaited plasma protein-A,free beta human chorionic gonadotropin, activin A and inhibin A at 22+0 to 24+6weeks gestation. Ultrasound Obstet Gynecol 2006;27(6):658-63.

14. Florio P, Ciarmela P, Luisi S, Palumbo MA, Lambert- Messerlian G, Severi FM et al. Pre eclampsia with fetal growth retardation: placental and serum activin a and inhibin A levels. Gynecol endocrinol 2002;16(5):365-72.

15. Fitzgerald B, Levytska K, Kingdom J, Walker M, Baczyk D, Keating S. Villous trophoblast abnormalities in extremely preterm deliveries with elevated second trimester maternal serum hCG or inhibin A. Placenta 2011;32(4):339-45.

16. Dayal M , Gupta P , Varma M , Ghosh UK , Bhargava A. Role of Second Trimester Maternal Serum Markers as Predictor of Preeclampsia. J Obstet and Gynecol Ind. 2011 pg 38-41.

17. Stewart CEH, Rotwein P. Growth, differentiation, and survival: multiple physiological functions for insulin like growth factors./ Physiol Rev 1996;76:1005-1026.

18. Hernandez- Valencia M, Zarate A, Ochoa R, Fonseca ME, Amato D, De Jesus Ortiz M. Insulin like growth factor-1, epidermal growth factor and transforming growth factor beta expression and their association with intrauterine fetal growth retardation such as development during huma pregnancy. Diabetes Obes Metab 2001;3(6):457-62.

19. Giudice LC, Martina NA, Crystal RA, et al. Insulin like growth factor binding protein-1 at the maternal- fetal interface and insulin like growth factor-1, insulin like growth factor II and insulin like growth factor binding protein-I in the circulation of women with severe preeclampsia. Am J Obstet Gynecol 1997;176:751-8.

20. Halhali A, Tovar AR, Torres N, Bourges H, Garabedian M, Larrea F. Preeclampsia is associated with low circulating levels of insulin – like growth factor-1 and 1,25 dihydroxyvitamin D in maternal and umbilical cord compartments. J Clin Endocrinol Metab 2000;85:1828-33.

21. Shang LX, Wang J, Zhang LJ, Gao H, Qu DY, Wang JH. Relationship between changes of insulin like growth factor-1 and insulin like growth factor binding protein -1 in maternal serum and placenta and pathogenesis of hypertensive disorder complicating pregnancy. Zhonghua Fu Chan Ke Za Zhi 2005;40(8):255-9.

22. Chiesa C, Osborn JF, Haass C, Natale F, Spinelli M, Scapillati E et al. Ghrelin, leptin, IGF-1, IGFBP-3, and insulin concentrations at birth: Is there a relationship with fetal growth and neonatal anthropometry? Clin Chem 2008;54(#):550-8.

23. Lindsay RS, Westgate A, Beattie J, Pattison NS, Gamble G, Mildenhall LFJ et al. Inverse changes in fetal insulin-like growth factor (IGF)-1 and IGF binding protein-1 in association with higher birth weight in maternal diabetes. Clin Endocrinol 2007;66(3):322-328.