PRIMARY AMENORRHOEA - AN ANALYSIS OF 5 CASES

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SUMMARY

Fifty-two adolescents with primary anenorrhoea were thoroughly investigated. Turner's syndrome and Rokitansky Kuster Hauser syndrome were the commonest etiological factors in our series. 5.7% of the patients were found to have genital Koch's. Associated cardiac and renal anomalies were frequently encountered. Ultrasonography was found to be a useful non-invasive diagnostic tool in these patients.

Introduction

The hallmark of the adolescent years is the initiation and completion of the pubertal process. Menarche and the initiation of regular menstrual cycles signal an uneventful completion of the pubertal process.

The development of normal menstrual function is documentation that the neuroendocrine, gonadal and anatomic components of the reproductive system are intact and mature. Amenorrhoea may be the first obvious sign of an abnormality in both reproductive and nonreproductive systems.

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Materials and methods

A total number of 52 adolescents coming to the out-patient department with complaints of primary amenorrhoea were screened.

After clinical examination, these cases were thoroughly investigated. T3, T4 levels, FSH, LH and prolaction levels, 17KS and DHEAS levels were estimated whenever necessary. X-ray skull and ultrasonography were done.

Intravenous pyelography and 2-dimensional echocardiography were performed to look for associated congenital anomalies.

Karyotyping was done on the peripheral blood of all patients with elevated gonadotrophin levels. Buccal smear was studied for X and Y chromosome. Diagnostic laparoscopy was done in indicated cases.

AN ANALYSIS OF 52 CASES

Results and discussion

(Insert Table I here)

TABLE I

Age distribution at presentation with primary amenorrhoea

Age in years	No. of case	No. of cases		
Less than 14	Contract and a second	2		
14-16	2	6		
16-18	1	2		
18-20		2		

Rokitansky Kauster Hauser syndrome was the commonest.

(Insert Table III here)

TABLE III

Endoscopy findings in vaginal agenesis and CIOF

	TFS RKHS CIOF		
No. of cases	1	5	6
Normal Uterus	YX IN 1		3
Hypoplastic or			
nodular uterus	in the second	5	3
Normal ovary	1	5	2
Streak ovary		-	4
Pelvic kidney	-	1	1

Majority of the cases presented between 14-16 years of age.

(Insert Table II here)

TABLE II

Etiologic factors of primary amenorrhoea

CCOF = chromosomally competent ovarian failure

20 15	CCOF 46XX 46 X Y	Irreversible		Rest	Convert Lower
10 5	CIOF 25% 45 XO 45 XO/XY 46	Reversible Physiologic delay 15%	att att att	Anatomic RKH 16%	
FSH/LH El	Hypergonadotrophic Hypogonadism	Hypogonadotropic Hypogonadism	Eu	gonadism	

Turner's syndrome was the most commonly encountered problem and was seen in 25% cases. Of the patients with eugonadism, Ultrasonography was performed on all cases. Some of them had an additional endoscopy

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done on them. Pelvic kidney was found in 1 case each of RKHS and CIOF.

(Insert Table IV here)

TABLE IV

Karyotype profile in chromosomally incompetent ovarian failure

Karyotype		No. of cases
Classical 45%	9	6
Y cell line 45 X/46 XY		2
Structural anomalies of X		3
X mosaic cell line		2
Total		13

Two patients showed a Y cell line mosaic. This is of special importance as a Y cell line especially if present in the gonads as well, can lead to future gonadal malignancies.

TABLE V

Primary amenorrhoea - associated anomalies

Etiology	Somatic	Cardiac	Renal	
CIOF	23%	23%	7.6%	
RKHS	2.5%	~	37.5%	
Others	16%	3.22%	6.4%	

37.5% of the patients with RKHS had renal anomaly of which lateral renal agenesis was the commonest. Coarctation of aorta and horseshoe kidney are frequently associated with imcompetent ovarian failure.

Conclusion

The commonest causes of primary amenorrhoea are chromosomally incompetent failure and anatomic defects. Contrary to the previously accepted idea that physiologic delay was the commonest cause of primary anenorrhoea, in our series it took a third place. In our series 5.7% of the patients with primary amenorrhoea had genital Koch's - hence in a developing country like India where Koch's is endemic, failure of an amenorrhoic patient to have withdrawal bleeding with estrogen and progesterone therapy should prompt one to suspect genital Koch's. This can be confirmed by histopathology, dilatation and curettage, laparoscopy and guinea pig inoculation.

The high incidence of renal and cardiac anomalies in CIOF and RKHS should prompt one to investigate all these patients with intravenous pyelography and echo-cardiography.

Ultrasonography and laparoscopy are very useful in investigating patients with primary amenorrhoca.

Ultrasound being simple, safe and nearly as predictive as endoscopy, it is now preferred.

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