Study of Primary Amenorrhea with etiology in out door patient of Rupganj Upazilla Health Complex

S Sultana¹, N Sultana², S A Ullah³, S A Walida⁴, P Akter⁵, F Ahmed⁶

Abstract

Background: Amenorrhea means without menstruation. Primary Amenorrhea is the absence of menstruation by 16 years of age in the presence of normal secondary sexual characteristics or by 14 years of age if secondary sexual characteristics have not developed.

Objectives: To explore the prevalence of Primary Amenorrhea with etiology in outdoor patient with limited investigation sources.

Methods: This was a cross sectional study conducted at Rupganj Upazila Health Complex (UHC) from January 2014 to June 2016. During this period total 2500 patients attended at Gynae out patient department (OPD). Among them 45 patients was sorted out with Primary amenorrhea. History and physical examination was done for presence or absence of secondary sexual characteristics, external genital abnormalities. Investigations were ultrasonogram (USG), Hormonal test were Follicle stimulating Hormone (FSH), Lutilizing hormones (LH), Serum Prolactin, Thyroid Stimulating Hormone. Karyotyping done. Data were analysed by using SPSS version 20.

Result: In our study prevalence of Primary Amenorrhea was 1.8%. Among them infantile uterus was 17.78%, imperforated hymen was 13.33%. Turner mosaic 11.4%, mullarian agenesis 6.66%, gonadal agenesis-4.45%, hypothyroid 8.89%, hyperprolactenemia-8.89%, and 11.11% patients were with positive progesterone challenge test without anatomical and hormonal abnormalities.

Conclusion: Considering the study result, Infantile uterus was the most prevalent etiological factor of amenorrhea followed by imperforated hymen and Turner mosaic.

Key word: Primary Amenorrhea, infantile uterus, imperforated hymen, Turner mosaic.

Introduction

Primary Amenorrhea is one of the common reproductive disorder affecting females. It leads to the absence of menstruation in the reproductive age group in females. Reported incidence of Primary Amenorrhea is 1-3% of women in reproductive age group. There are many causes which leads to Primary Amenorrhea including genetic aberrations. Various factors such as anatomical, genetic and hormonal factors reported to influence Primary Amenorrhea. The recommended evaluation for amenorrhea aimed to divide the reproductive system into components like genital outflow tract, uterus, the ovary, the pituitary and to assess the functional integrity of each, starting at the lowest level (the genital outflow tract) and upto the highest levels (hypothalamus).

The most common etiology of Primary Amenorrhea is- gonadal dysgenesis (40%), Hypothalamic Amenorrhea (30%). The genital examination is abnormal in 15% of patients with Primary Amenorrhea. In rare condition more than one component of hypothalamus- pituitary-ovary (HPO) axis and genital tract are affected. Other than each 20% hypoprolactenemia and 30% chance of weight related Amenorrhea.

The etiologic factors of primary amenorrhea

Group I- End organ failure/ outflow tract obstruction

a) Mullerian agenesis
b) Transverse vaginal septum
hypogonadotropic hypogonadism includes
(a) Congenital abnormalities
(b) Endocrine disorders
(c) Tumor
(d) Systemic illness (2.6%)
(e) Eating disorder (2.3%).

a) Congenital abnormalities that can cause hypogonadotropic hypogonadism include the following
Isolated GnRh deficiency (8.3%)
Forms of hypopituitarism (2.3%)
Congenital CNS defect (0.8%)
Constitutional delay (6%).

b) Endocrine disorders that can cause hypogonadotropic hypogonadism includes-
Congenital Adrenal Hyperplasia (CAH) (0.8%)
Cushing Syndrome (0.4%)
Hyperprolactinemia (1.9%)

Tuberculous endometritis
Male pseudo hermaphroditism- complete testicular feminization.

Group II- Gonadal failure
a) Gonadal dysgenesis (46XX, 45XO, Mosaic)
b) Agonadism
c) Post chemotherapy

Group III- Pituitary cause
a) Hyperprolactinemia
b) Prolactinoma

c)    Tuberculous endometritis
d)    Male pseudo hermaphroditism- complete testicular feminization.

Group IV
a) Hypogonadotropic hypogonadism
b) Hypothalamic dysfunction

Group V
a) Primary hypothyroidism
b) Congenital adrenal hyperplasia
c) Androgen secreting tumour

Secondary amenorrhea is absence of menstruation for their normal cycle or for six months.

Physiological Amenorrhea occurs before puberty, during pregnancy, lactational period and during menopause.

Pathological Amenorrhea are of 3 types-

- Amenorrhea without secondary sexual characteristics.
- Amenorrhea with secondary sexual characteristics and anatomic abnormalities.
- Amenorrhea with secondary sexual characteristics and nonanatomic cause.

For majority of pubertal girl’s menstruations is the final result of series of events which result in sexual maturity. Maturation of hypothalamus through several years of late childhood begins a cascade of events of the normal menstrual cycle and menstruation.

First and foremost, it is imperative to rule out pregnancy. Additional diagnosis of Primary Amenorrhea usually result from genetic or anatomic abnormality. The relative prevalence of Primary Amenorrhea includes hypergonadotropic hypogonadism (48.5%), hypogonadotropic hypogonadism (27.8%) and eugonadism (pubertal delay) with normal gonadotrophin (23.7%).

The hypergonadotropic hypogonadism category includes (a) patients with abnormal sex chromosome (i.e. turners syndrome) who make up 29.7% of all Primary Amenorrhea cause and those (b) Patient with normal sex chromosomes. The later group includes both patients who are 46xx(15.4%) and those 46xy(3.4%).

We conducted this study to evaluate the etiology of primary amenorrhea in women attending at a Primary health care centre in Bangladesh.

Methods
The study design was descriptive cross sectional and conducted at Rupganj Upazilla Health Complex (UHC) from 2014 January to June 2016. Total 2400 patients were attended at Gynae OPD among them 45 patients with primary amenorrhea were sorted out following nonprovability purposive sampling.

Then proper case history was taken and physical examination was done for presence or absence of secondary sexual characteristics, external genital abnormalities, and several laboratory investigations (ultrasonogram (USG), Hormonal test were Follicle stimulating Hormone (FSH), Lutilizing hormones (LH), Serum Prolactin, Thyroid Stimulating Hormone.
Karyotyping (done from Bangladesh Institute of Research and Rehabilitation in Diabetes, Endocrine and Metabolic Disorder (BIRDEM) and Bangabandhu Sheikh Mujib Medical University (BSMMU)). Hormonal assay done to sort out endocrine disorder. Ultrasonogram (USG) was done to find out any abnormalities of internal genital organs-infantile uterus, Mullerian agenesis, gonadal agenesis, imperforated hymen etc. Karyotyping was done to find out chromosomal anomaly.

Those patients who were within normal BMI and presence of secondary sexual characteristics and normal biochemical findings were given progesterone challenge test.

Data collected from individual history and data sheet. Prior to data collection ethical clearance was taken from authority and informed consent was taken from patients. Confidentiality and privacy was maintained while data collection was done. Data were cleaned, edited and analyzed by using SPSS version 20. The frequency was calculated and percentage was done to find out the prevalence of etiology.

**Result**

Among 45 patients 43 patients (95.55%) was found with secondary sexual characteristics (Breast development, Pubic and axillary hair and with normal external genitalia). 2 (4.45%) patients found without secondary sexual characteristics but with normal external genitalia. (Table-1)

<table>
<thead>
<tr>
<th>Secondary Sexual Characteristic</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Present</td>
<td>43</td>
<td>95.55%</td>
</tr>
<tr>
<td>Absent</td>
<td>02</td>
<td>04.45%</td>
</tr>
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</table>

USG showed that Infantile uterus is about <5 cm in length but gonad present in 17.78%, Uterus and Adenexa present but haematocolpas, haematometra that is imperforated hymen 6 (13.33) patients out of 45 patients. About 5(11.11%) patients are found with streak gonad/complex gonadal structure. Gonad present but uterus absent i.e Mullerian agenesis 3(6.66%). But gonad absent uterus present in 2(4.45%) patients. About 21(46.67%) patients are found with normal uterus and normal gonad on USG finding. (Table-II)

<table>
<thead>
<tr>
<th>USG finding</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal internal genitalia with imperforated hymen</td>
<td>06</td>
<td>13.33%</td>
</tr>
<tr>
<td>Infantile uterus</td>
<td>08</td>
<td>17.78%</td>
</tr>
<tr>
<td>Mullerian agenesis</td>
<td>03</td>
<td>06.66%</td>
</tr>
<tr>
<td>Gonadal agenesis</td>
<td>02</td>
<td>04.45%</td>
</tr>
<tr>
<td>Streak/complex gonadal structure</td>
<td>05</td>
<td>11.11%</td>
</tr>
<tr>
<td>Normal uterus with normal gonad</td>
<td>21</td>
<td>46.67%</td>
</tr>
</tbody>
</table>

Result on hormonal analysis showed that 22 (48.89%) had normal level. 5 patients had increased patients FSH, LH increased in 10 (22.22%) patients, increased TSH in 4 (08.89%) patients, increased serum Prolactin level in 4 (08.89%) patients. (Table III)

<table>
<thead>
<tr>
<th>Hormonal Analysis</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Within Normal</td>
<td>22</td>
<td>48.89%</td>
</tr>
<tr>
<td>FSH-- increased</td>
<td>05</td>
<td>11.11%</td>
</tr>
<tr>
<td>LH-- increased</td>
<td>10</td>
<td>22.22%</td>
</tr>
<tr>
<td>TSH-- increased</td>
<td>04</td>
<td>08.89%</td>
</tr>
<tr>
<td>S.Prolactin-- increased</td>
<td>04</td>
<td>08.89%</td>
</tr>
</tbody>
</table>

Turner mosaic 45XO found in 7(38.89%) and 11(61.11%) patients with 46XX karyotype (Karyotyping done in 18 patients) (Table IV)

<table>
<thead>
<tr>
<th>Karyotype</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>45XO-(Turner’s mosaic)</td>
<td>07 patients</td>
<td>(38.89%)</td>
</tr>
<tr>
<td>46XX-</td>
<td>11 patients</td>
<td>(61.11%)</td>
</tr>
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Among 45 patients of Primary Amenorrhea those with all hormonal, USG finding were normal, progesterone challenge test was positive. They had normal menstrual flow after withdrawing of 7 days progesterone orally.

All these patients 45 family history was negative for any gynecological issue.

**Discussion**

At Rupganj UHC from January 2014 to June 2016 about 2500 patients attended at Gynae OPD. Among them 45 patients was sorted out with Primary amenorrhea prevalence was 1.8%.

In our study infantile uterus was 17.77%, Mullerian agenesis-6.67%, gonadal agenesis-4.45%. Study of Primary Amenorrhea by Elisabeth H. Quint, M.D. and Yolanda R. Smith MD., MS showed similar result. On USG examination their study revealed infantile uterus with normal ovaries and FSH and Estradiol levels were normal. 7

In our study on ultrasonogram finding revealed that uterus and gonad present but haematocolpos and haematometra with imperforated hymen found in 13.33% and complex gonad/streak gonad in 11.11% cases. On the other hand Firouzeh Ghaffari, Fatemeh Keikha and Arezoo Arabipoor reported Primary Amenorrhea with two etiologies–Hypothalamic Amenorrhea and transverse vaginal septum with no haematocolpos. 3

The current study revealed hormonal analysis of FSH, LH, Prolactin, TSH were normal in 48% cases and rest cases showed increased level of different hormones. Similar result was found in study done by Z. Nazir et al. 10

In our study, karyotyping was done in 18 patients. Among them
7(38.89%) patients were 45XO and 11(61.11%) patients were 46XX. In a study by Amdhu et al showed that 305 cases were confirmed to have chromosomal abnormalities. Among 45 patients of Primary Amenorrhea 5(11.11%) with all hormonal, USG finding normal and progesterone challenge test was positive, that is, they have normal menstrual flow after withdrawing of 7 days progesterone orally. All of the patients (45) family history was negative for any gynaecological issues. A study by M. Behera was also found similar result.9

**Conclusion**

Considering the study result, Infantile uterus was the most prevalent etiological factor leading to amenorrhoea followed by imperforated hymen and Turner mosaic.

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