Thai Girl with Ring Chromosome 18 (46XX, r18)

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Chromosomal anomalies occur in 0.4% of live births. Ring chromosomes have been found for all human chromosomes and when it is replacing a normal chromosome, it results as a partial monosomy. The phenotype often overlaps that seen in comparable deletion syndromes of the same chromosomes. In the present report the authors describe the clinical manifestations of a girl with ring chromosome 18 (46XX,r18) including dysmorphic features, failure to thrive, global delay of development, hypothyroidism, atopic dermatitis, bilateral chronic otitis media, aortic regurgitation with patent foramen ovale and immunoglobulin A deficiency.

Keywords: Ring chromosome 18, Thailand, Pediatric, IgA deficiency, Mental retardation

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Chromosomal anomalies occur in 0.4% of live births(1). They are an important cause of mental retardation and congenital anomalies. Chromosome anomalies are found in number and structure. Abnormalities of chromosome structure include deletions, translocations, inversions, ring chromosomes, duplications and insertions. Formation of a ring chromosome involves a deletion at each end of the chromosome then the “sticky” ends join to form a ring. Though it is rare, ring chromosomes have been found for all human chromosomes. When it replaces a normal chromosome, it results in a partial monosomy. The phenotype often overlaps that seen in comparable deletion syndromes of the same chromosomes. In the present report, the authors to describe the clinical manifestations of a girl with ring chromosome 18.

Case Report

A 2 year old girl presented with a history of chronic eczema and chronic otitis media for 1 year. She had failure to thrive and delayed developmental milestones. Physical examination revealed an active dysmorphic child with light brown hair. Dysmorphic features included a flattened forehead; deep-set eyes, hypotelorism, bilateral ptosis, low set ears, high arch palate, protrusion of the maxilla and micrognathia. Her height, weight and head circumference, 65 cm, 5.3 kg and 39 cm were all less than the 3rd percentile. The anterior fontanel was 1.5 x 1.5 cm, the posterior fontanel had closed. Purulent discharge was seen in both ear canals. Her nipples were widely spaced. She had a systolic ejection murmur of grade III/VI at her left upper parasternal boarder. On abdominal examination, mild hepatomegaly and an umbilical hernia were noted. She had ulnar deviation of hands, genu valgum and over riding of 2nd finger over 1st toes in both feet. Her skin was dry with fine scales at the fore head, erythematous papules with crust and serum oozing and scales at the flexor area of all extremities.

She was a near term baby delivered by normal labor at 36 weeks of gestational age. The birth weight was 2100 gm. During the neonatal period she required admission due to jaundice and poor feeding, she received 4 days of phototherapy and was fed via orogastric tube feeding during and after discharge. The total admission period was 2 months. She was the third child of a 38 year old mother and 26 year old father; their first 2 children 11-year-old son and 9-year-old daughter were all healthy.
Blood samples revealed fasting blood sugar; 72 mg/dl, TSH; 0.05 (0.4-3.1 mIU/ml), FT4; 0.4 (0.86-1.87 ng/dl). Her serum Immunoglobulin levels were; IgG 2407.77 (1244-1888), IgM 186.79 (104-202), IgA < 40.69 (80-240). At the age of 2 years old, her bone age corresponded with the age of 6 months. Mild hepatomegaly and bilateral renal parenchymal disease were seen on abdominal ultrasonography. On cardiologic evaluation, sinus tachycardia with heart rate of 160 beats per minute and QRS axis of 240° were present on electrocardiography. Echocardiography findings included patent foramen ovale 0.36 cm in diameter with left to right flow, aortic valve regurgitation with pressure gradient 13.4 mmHg and EF 59.4%. She had global delay of development. Despite her age, her average abilities were equal to a 4 month old child. The personal-social aspect was equal to 5 months, fine motor 4 months, language 3 months and gross motor 5 months. Karyotype analysis using G-banding technique (Fig. 1) resulted as 46XX, r18.

The diagnosis of ring chromosome 18 was made, with failure to thrive, global delay of development, hypothyroidism, atopic dermatitis, bilateral chronic otitis media, aortic regurgitation with patent foramen ovale, and immunoglobulin A deficiency. After admission and further workup, she was treated with antibiotics for chronic otitis media, the pus cultures were positive for *P. aeruginosa*. Thyroid hormone was supplemented. Her skin lesions improved after treatment with topical corticosteroids.

**Discussion**

There are no reports on clinical manifestations of Thai patients with ring chromosome 18. The

![Fig 1. Patient's Karyotype analysis by G-banding technique, the arrow indicates the ring chromosome](image-url)
authors describe the clinical manifestations of a Thai girl with ring chromosome 18 (46XX, r18). The syndrome of ring chromosome 18 results from deletions at both ends of the chromosome, then the joining of the ends of the p and q arms back together. There is chromosome material lost from both arms. Individuals with ring chromosome 18 have a combination of 18p deletion and 18q deletion (2-7). This corresponds with the parts of the chromosome that are deleted. All features of this syndrome have been described in someone with 18p deletion or 18q deletion (8). This patient shares clinical manifestations of both chromosome 18p and 18q deletion syndrome (Tables 1(9) and 2 (10)), though correlated with chromosome 18q deletion syndrome more than chromosome 18p deletion syndrome.

Hypothyroidism has been reported in ring chromosome 18 syndrome (3, 11, 12). The presented patient had delayed development and delayed bone age that corresponded with secondary hypothyroidism and ring chromosome 18 syndrome. Growth hormone deficiency, also reported in this syndrome (13, 14), could be another cause of the failure to thrive. However, in the presented patient growth hormone studies were not done due to economic causes. Other autoimmune endocrinopathies reported with this syndrome, such as insulin dependent diabetes mellitus (13) was not detected in the presented patient. Immunoglobulin A deficiency which is frequently associated with ring chromosome 18 syndrome (4, 15) was also noted. This should be kept in mind, especially when considering the use of blood products in this patient group. Close follow up is indicated. Further cytogenetic studies, DNA analysis and Fluorescence In Situ Hybridization studies should be done to identify location of chromosomal breakpoints.

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References

9. United States National Library of Medicine and

Table 1. Clinical features of chromosome 18p deletion syndrome in this patient

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<thead>
<tr>
<th>Head and neck</th>
<th>microcephaly, brachycephaly, prognathism, midfacial hypoplasia</th>
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<tr>
<td>Mouth and oral structures</td>
<td>highly arched palate</td>
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<td>Skin</td>
<td>eczema</td>
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<td>Cardiovascular system</td>
<td>atrial and auricular septal defect</td>
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<td>Immunologic system</td>
<td>immunoglobulin A deficiency</td>
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<td>Growth and development</td>
<td>growth and mental retardation</td>
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Table 2. Clinical features of chromosome 18q deletion syndrome in this patient

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<th>Head and neck</th>
<th>microcephaly, brachycephaly, prognathism, midfacial hypoplasia</th>
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<tbody>
<tr>
<td>Eyes</td>
<td>deeply set eyes and hypotelorism</td>
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<tr>
<td>Mouth and oral structures</td>
<td>highly arched palate</td>
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<tr>
<td>Hand and foot</td>
<td>abnormal toes</td>
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<tr>
<td>Immunologic system</td>
<td>immunoglobulin A deficiency</td>
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<tr>
<td>Growth and development</td>
<td>growth, speech, and mental retardation</td>
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รายงานผู้ป่วย; เด็กหญิงไทยที่มีโครโมโซมแท่งที่ 18 เป็นโครโมโซมวงแหวน 1 แท่ง (46XX, r18)

รสรวมค์ ศรีพานิชกุลชัย, ชายภูร่ย์ ศุภคุณภิญโญ, จรูญ เจตศรีสุภาพ, สมาน เลืองวัฒนาวิช

ความผิดปกติของโครโมโซมพบได้ 0.4% ของขั้นตอนการเกิดมีริ้ว การเกิดโครโมโซมวงแหวนพบได้ในทุกโครโมโซมของมนุษย์ การเกิดโครโมโซมวงแหวนพบที่โครโมโซมปกติ 1 แท่ง ผู้ป่วยจะมีการเจริญถดถอยเนื่องจากความผิดปกติของโครโมโซมที่หายไป รายงานฉบับนี้ บรรยายลักษณะของผู้ป่วยเด็กหญิง อายุ 2 ปีที่มีโครโมโซมวงแหวน 18 เป็นโครโมโซมวงแหวน 1 แท่ง (46XX, r18) โดยเกิดลักษณะเลี้ยงไม่โต ภาวะขาดธัยรอยด์ฮอร์โมน ผื่นผิวหนังอักเสบ หูชั้นกลางอักเสบเรื้อรัง หูชั้นกลางรั่ว ลิ้นหัวใจเอออร์ติกรั่ว ผนังกั้นหัวใจ ห้องบนรั่ว (patent foramen ovale) และมีระดับของ อิมมูโนโกลบูลินเอ ต่ำ