Park syndrome

**Eponyms:**
- Baird-Lowry syndrome
- Klippel-Feil deformity-deafness-absent vagina

**Inheritance:**
supposed autosomal dominant

**Semeiological Synthesis:**

**Group Sub group Signs:**

<table>
<thead>
<tr>
<th>AUDITORY DISORDERS</th>
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</thead>
<tbody>
<tr>
<td>deafness</td>
</tr>
<tr>
<td>deafness conductive type</td>
</tr>
<tr>
<td>deafness sensorineural, including unspecified type</td>
</tr>
<tr>
<td>deafness, in syndromic association</td>
</tr>
<tr>
<td>ear, ossicular malformation</td>
</tr>
</tbody>
</table>

**GENITAL DISORDERS**

*female genitalia, modified not including ambiguity*

- vagina, atresia, absence

**genital dysfunctions**

- amenorrhea, primary

**uterus, anomalies**

- mullerian duct, agenesis/defects
- uterus, absent

**OROCRANIOFACIAL ANOMALIES**

*facies, modified appearance*

- facies, asymmetric, unilateral atrophy, hemifacial atrophy

**neck, modified appearance**

- pterygium colli, webbed neck, including: thickened nuchal fold, nuchal thickening

**OTHERS**

*supergroups*

- facio-genito-skeleto-urinary
- genito-skeletal disorders
- oto-skeleto-urological disorders

**PRENATAL-NEONATAL MODIFIED DATA**

*foetal changes*

- foetal changes, recognized by ultrasound techniques

**prenatal diagnosis**

- prenatal diagnosis, echographic

**SKELETAL DISORDERS**

*spine, changes*

- vertebra, fusion

**stature, growth, modified habitus**

- stature, short, including micromelia, including short limbs

**vertebral changes**

- vertebra, cleft, hemivertebra, including iniencephaly
- vertebral anomalies, unspecified type

**UROLOGICAL DISORDERS**

*kidney, malformations*

- kidney, ectopic

**Super group:**

- facio-genito-skeleto-urological disorders
- genito-skeletal disorders
- oto-skeleto-urological disorders
Super agg.  DEAFNESS
Aggregations:  deafness, in syndromic association

FOETAL CHANGES
foetal changes, recognized by ultrasound techniques

OTHER
mullerian duct, agenesis/defects

Differential diagnosis:
10050  Fragoso syndrome
14000  Klippel-Feil I syndrome
14001  Klippel-Feil II syndrome
14002  Klippel-Feil III syndrome
14003  Klippel-Feil IV syndrome
27545  Lampert syndrome
23715  Shaver syndrome

Bibliography  OMIM ID: 148860