The genetic aetiology of Silver–Russell syndrome

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ABSTRACT
Silver–Russell syndrome (SRS MIM180860) is a disorder characterised by intrauterine and/or postnatal growth restriction and typical facies. However, the clinical picture is extremely diverse due to numerous diagnostic features reflecting a heterogeneous genetic disorder. The mode of inheritance is variable with sporadic cases also being described. Maternal uniparental disomy (mUPD) of chromosome 7 accounts for 10% of SRS cases and many candidate imprinted genes on 7 have been investigated. Chromosome 11 has moved to the forefront as the key chromosome in the aetiology, with reports of methylation defects in the H19 imprinted domain associated with the phenotype in 35–65% of SRS patients. Methylation aberrations have been described in a number of other imprinted growth related disorders such as Beckwith–Wiedmann syndrome. This review discusses these recent developments as well as the previous work on chromosome 7. Other candidate genes/chromosomal regions previously investigated are tabled.

GENETIC AETIOLOGY
SRS is a clinically and genetically heterogeneous disorder. Autosomal dominant, autosomal recessive and X-linked inheritance models have all been reported.7 Both concordant8 and discordant9,10 monozygotic twins for SRS have also been found. Given this complex genetic aetiology, it is not surprising that a number of different chromosomes and potential candidate genes have been implicated in the cause of SRS. Up to 2005, the most widely discussed chromosome involvement was with chromosome 7, but recent findings have shown that loci on chromosome 11 have a big part to play in the origins of this disorder. The implications of these exciting new findings are discussed in detail below.

Chromosome 7
Chromosome 7 as a candidate locus for SRS has been studied in detail after maternal uniparental disomy (mUPD) 7 was found in a subset of patients with pre- and postnatal growth restriction.11–14 It was postulated that the growth restriction might be due to growth related imprinted genes located on this chromosome.

Imprinted genes represent a very small proportion of the total gene complement, which differ from the rest in the manner of their transcriptional regulation. Unlike most genes that are active from both maternally and paternally inherited chromosomes (that is, biallelic), imprinted genes are only active from one parental allele (that is, monoallelic). For germ line imprinted genes, this is dictated in the egg or sperm. Imprinted genes are therefore regulated according to parent-of-origin, and function in a hemizygous state. An imprinted gene may be paternally expressed and maternally imprinted (silenced) or maternally expressed and paternally imprinted (silenced).

This hypothesis was elegantly demonstrated by pioneering work in the 1980s by Barton et al15 and McGrath and Solter16 who created diploid mouse embryos with either gynogenetic (completely maternal) or androgenetic (completely paternal) genomes to examine the phenotypic effect on conceptuses. Gynogenetic conceptuses produced only embryonic material with little or no placenta whereas androgenetic conceptuses displayed poor embryonic development with recognisable trophoblast tissue and yolk sacs. It was obvious from these experiments that the maternal and paternal genomes made differential contributions to fetal growth and development. This hypothesis was further defined by the creation of mouse strains with Robertsonian and reciprocal translocations to create artificial UPDs for sub-chromosomal regions. Interestingly, some chromosomes in the UPD state had no visible phenotypes whereas
others showed profound growth related effects. One of the many cited examples is that of mice with maternal or paternal UPD for part of mouse chromosome 11, which is syntenic with human chromosome 7. Mice with mUPD11 were on average 30% smaller than their wild type littermates, independent of sex, while the pUPD11 mice were 30% larger than their wild type littermates. This demonstrated a polarising epigenetic effect on the same phenotype. It is now accepted that, in general, paternally expressed genes show promotion of fetal growth whereas maternally expressed genes have a tendency to restrict fetal growth. This imprint, which effectively silences one parental copy of a gene, is epigenetic, and is reset during gametogenesis.

The findings that maternal UPD7 in humans is associated with pre- and postnatal growth restriction, and that imprinted genes have a phenotypic dosage effect in mice, suggested chromosome 7 as a possible candidate for SRS pathogenesis. Several groups therefore screened cohorts of clinically defined SRS patients both for mUPD7 and/or imprinted gene(s) on chromosome 7 that may contribute to the observed growth disorder. UPD7 is readily detectable using polymorphic short tandem repeat (STR) microsatellite markers by polymerase chain reaction (PCR) or variable number of tandem repeat (VNTRs) by Southern blotting. Of the 98 SRS patients collectively screened by these groups, nine were found to have mUPD7 and it is now widely acknowledged that this mechanism is responsible for approximately 10% of SRS cases. As an alternative to disrupting imprinted gene expression, the hypothesis that the UPD might be uncovering a recessive allele was also tested. Several examples of this have been described for chromosome 7 where patients with cystic fibrosis appear to be homozygous for delta508 despite only having one carrier parent. This turns out to be due to isodisomy, where the patient inherits two copies of the same chromosome from one parent while the normal chromosome is excluded. Freece et al therefore analysed five mUPD7 patients for common regional isodisomic inheritance using 40 polymorphic markers distributed along the length of chromosome 7 with an average distance between markers of <10 cM. They found no common region of isodisomy among the five probands, suggesting the phenotypic effect of chromosome 7 is either due to exposure of recessive alleles for several different genes or much more likely due to a common imprinting effect. The observation of mUPD7 in SRS as a genetic aetiology appears to be an almost exclusive phenomenon as it has not yet been reported in screening of individuals with intrauterine growth restriction (IUGR) or pre- and/or postnatal growth restriction.

Interestingly, Hoglund et al and Pan et al describe single case reports each of complete paternal isodisomy for chromosome 7 with normal growth phenotype. This contrasts sharply with the overgrowth phenotype seen in the syntenic pUPD11 mice and suggests that the human phenotype is related to maternal growth restricting genes rather than paternal growth promoting genes. However, a single case report of a boy with cystic fibrosis with complete paternal isodisomy for chromosome 7 displaying overgrowth was recently published, reopening the possibility of paternally active imprinted genes on this chromosome playing a role in growth. Two key candidate regions on chromosome 7 have been delineated as the most likely areas to contain the genes that may be causative factors in SRS.

7q32

Kobayashi et al suggested the human equivalent of the imprinting mouse Peg1/Mest (mesoderm expressed transcript) gene as a candidate for SRS. They developed a novel subtraction–hybridisation method to isolate imprinted genes systematically. Of eight paternally expressed genes (Pegs), Peg1/Mest was identified as the most abundantly expressed in embryos, specifically in the mesoderm. Mest maps to mouse proximal chromosome 6, which shares conserved synteny with human chromosome 7q21-qter. This region contains the paternally expressed MEST gene, which was the first imprinted gene to be identified on chromosome 7. Based on its imprinting status and the mouse phenotype, MEST has been investigated extensively as a candidate for SRS, but sequencing and methylation studies have as yet found no evidence for an involvement.

An SRS patient with a rare segmental mUPD for the region 7q31-qter further implicated this region to be harbouring a potential candidate gene(s) for SRS. Recently, a case report of a girl born after assisted reproductive technology (ART) and diagnosed with SRS was investigated for methylation status of the MEST region. They observed partial hypermethylation of the MEST DMR and speculated that this may have some effect on this paternally expressed growth promoting gene.

Several other imprinted genes were found in the vicinity of MEST, including the paternally expressed intronic transcript

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**Box 1: Characteristics of Silver–Russell syndrome (SRS)**

**Physical features**
- Facial dysmorphism: classic facial features of small triangular facies, high forehead with pointed chin and down turned corners of the mouth
- Short stature persisting into adulthood
- Skeletal asymmetry
  - Preservation of occipitofrontal circumference
  - Clinodactyly
  - Camptodactyly ± distal arthrogyrosis
  - Syndactyly of second and third toes
  - Hemihypertrophy
- Urogenital dysmorphism:
  - Hypospadias
  - Posterior urethral valves
  - Inguinal hernia

**Clinical features**
- Birth weight \( \leq -2 \) SD from the mean. Usually followed by failure to thrive
- Poor head control during infancy due to relatively large head compared to body as well as general motor dysfunction due to lack of muscle bulk and strength
- Feeding difficulties in infancy and childhood leading to fasting hypoglycaemia
- Increased sweating during infancy
- Gastrointestinal symptoms including gastro-oesophageal reflux, oesophagitis and food aversion
- Developmental delay which may be associated with mental compromise in a subset of patients
- Delayed bone age
- Cardiac defects
- Malignancies


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References:
19 20 Several examples of this have been described for chromosome 7 where patients with cystic fibrosis appear to be homozygous for delta508 despite only having one carrier parent. This turns out to be due to isodisomy, where the patient inherits two copies of the same chromosome from one parent while the normal chromosome is excluded. Freece et al therefore analysed five mUPD7 patients for common regional isodisomic inheritance using 40 polymorphic markers distributed along the length of chromosome 7 with an average distance between markers of <10 cM. They found no common region of isodisomy among the five probands, suggesting the phenotypic effect of chromosome 7 is either due to exposure of recessive alleles for several different genes or much more likely due to a common imprinting effect. The observation of mUPD7 in SRS as a genetic aetiology appears to be an almost exclusive phenomenon as it has not yet been reported in screening of individuals with intrauterine growth restriction (IUGR) or pre- and/or postnatal growth restriction.27

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Several other imprinted genes were found in the vicinity of MEST, including the paternally expressed intronic transcript
from the biallelically expressed gamma-2 COP (nonclathrin-coat-protein) (COPG2/T1), paternally expressed MEST/T1 (also known as PEG1/AS) and the maternally expressed carboxypeptidase A4 (CPA4). CPA4 has a role in cell proliferation and differentiation. However, no mutations were seen for CPA4 in the 10 SRS patients screened or in other imprinted genes in this region. Neighbouring genes flanking these imprinted genes were all shown to be biallelic, delineating the extent of a small imprinting cluster. Meanwhile, mice either targeted for mutation of the homologous Mest gene or mice with mUPD for chromosome 6 (which are equivalent to null for the paternally expressed Mest gene), result not only in IUGR and reduction in postnatal survival but also in abnormal maternal behaviour including impaired placentaphagia. The latter finding provided the first evidence that normal and abnormal imprinting may also play an important role in behaviour.

7p11.2-p13
Joyce et al. reported a mother and daughter with some but not all of the key features of SRS. Cytogenetic and fluorescent in situ hybridisation (FISH) established that both mother and daughter had an inverted duplication of the proximal short arm of chromosome 7 encompassing the region 7p13-p12.1. This region harbours the growth factor receptor binding protein-10 (GRB10) gene which is imprinted in an isoform and tissue specific manner. GRB10 encodes a cytoplasmic adaptor protein which may facilitate linkage of cell surface tyrosine kinase receptors, such as the insulin receptor (IR) and insulin-like growth factor-1 receptor (IGFIR) to an uncharacterised mitogenic signalling pathway, and also binds epidermal growth factor receptor (EGFR) and the platelet derived growth factor receptor.

The orthologous Grb10 gene is located on mouse chromosome 11 which is associated with the growth imprinting phenotype described by Cattanach and Kirk. The GRB10 gene could be considered a very strong candidate for SRS. Monk et al. identified a maternally derived de novo duplication in a female patient with some of the classic features of SRS. The duplication involved 7p13-p11.2, encompassing GRB10 at 7p12.2 and the biallelically expressed insulin-like growth factor binding proteins 1 and 3 (IGFBP1 and IGFBP3). In addition to this family, a further two maternally inherited duplications encompassing GRB10, and several chromosomal disruptions in the 7p11.2-p13 region, have also been described. More recently a maternally inherited duplication of 7p12-p13 in a patient with cognitive deficit without features of SRS has been reported in the literature. This duplication does not extend to include GRB10 and delineates an SRS critical interval of 2.2 Mb containing only GRB10 and the biallelically expressed cordon bleu gene (COBL). Therefore the growth restriction observed is most likely due to the over-expression of the imprinted GRB10 gene.

The GRB10 gene is imprinted in a highly isoform and tissue specific manner. In both mouse and human, expression in the brain is derived from the paternal allele. In the mouse all other isoforms in embryonic and placental tissues are expressed from the maternally controlling this maternal allele. This maternal expression is not completely conserved in the human, with only the γ1 and γ2 isoforms imprinted in muscle, and expression in other tissues reported, derived from both parental alleles. Mice carrying maternally inherited targeted deletions of Grb10 show a growth phenotype identical to the phenotypes described for the pUPD11 and PatP11 mice, consistent with an involvement in SRS. In total, 76 SRS patients were screened for mutations in GRB10 but no pathogenic changes were found. Arnaud et al. and Monk et al. found no significant changes in the differentially methylated region (DMR) for GRB10 in 24 and 22 SRS patients, respectively. Therefore, despite the location and proposed function of GRB10 and its function in mice, there have not been any conclusive reports of SRS being caused by over-expression or loss of imprinting of GRB10, or by gain of function coding mutations.

Chromosome 11
The 11p15.5 region contains two clusters of imprinted genes each regulated by its own imprinting control region (ICR). H19 and insulin-like growth factor 2 (IGF2) are controlled by the H19 differentially methylated domain (DMR), which is the telomeric ICR1, while the genes within the KCNK1 domain are regulated by the centromeric KvDMR1 called ICR2. This region has become a key focus in the aetiology of SRS ever since the report of hypomethylation of the ICR1 in five of nine SRS patients studied by Gicquel et al. They hypothesised that epigenetic dysregulation of genes in the human 11p15.5 region encompassing the telomeric ICR1 region and the more centromeric ICR2 region involved in the aetiology of Beckwith–Wiedemann syndrome (BWS) (OMIM 150650) could be genotypically and phenotypically opposite for SRS.

BWS is characterised by macroglossia, pre- and/or postnatal growth greater than the 90th centile and abdominal wall defects. In addition, umbilical abnormalities and characteristic linear indentation of the earlobe are included in the diagnostic criteria. BWS patients are also at increased risk of tumours. Interestingly, these patients have large placentae that are almost twice the normal weight.

The genetic aetiology of BWS is extremely heterogeneous with most reported cases being sporadic. In a study of 106 sporadic BWS cases, 17% were due to paternal isodisomy of the 11p15 region which contains the reciprocally imprinted genes IGF2 and H19 (ICR1) as well as the KCNK1 domain (ICR2). As IGF2 is a potent prenatal growth factor, over-expression of IGF2 should lead to tissue hyperplasia. Weksberg et al. examined skin fibroblasts in five monozygotic twin pairs discordant for BWS and found loss of methylation of the maternal allele of the ICR2 in the affected twin only. Imprinting defects of the ICR2 that result in loss of expression of the cyclin dependent kinase inhibitor 1C (CDKN1C) gene are the most common epimutation found in 50% of BWS patients. Maternally inherited coding mutations of CDKN1C have also been reported in 5% of BWS patients. However, 10% of BWS patients exhibit hypermethylation of ICR1 and the promoter region, resulting in loss of expression of the non-coding H19 gene and the biallelic expression of IGF2.

Research into the imprinting control of this domain reveals a complicated regulatory mechanism. The ICR1 is a paternally methylated germline DMR, located 2–4 kb upstream of the H19 transcript that contains several CCCTC binding factor (CTCF) binding sites. The binding of CTCF confers ICR1 function as a methylation sensitive insulator between the multiple IGF2 promoters and enhancers located downstream of H19. On the unmethylated maternal allele, CTCF binds to form a boundary that prevents the IGF2 promoters interacting with the enhancers, whereas on the methylated paternal allele, CTCF cannot bind and the IGF2 promoters can freely associate with the enhancer to bring about expression from the paternal allele only. Therefore any gain or loss of methylation of ICR1 will result in aberrant IGF2 expression, potentially corresponding to the opposite growth phenotypes of SRS and BWS (fig 1).
lymphoblasts and fibroblasts of five SRS patients. The change was proposed to cause loss of methylation of the H19 promoter leading to biallelic H19 expression, downregulation of IGF2 and hence growth restriction. The degree of hypomethylation was correlated with the severity of the phenotype. Of critical note, one of the five SRS patients is a monozygotic twin discordant for SRS. Although her non-SRS twin showed the same partial hypomethylation in her leucocytes with concomitant biallelic expression of H19, skin fibroblast from the non-SRS twin showed no methylation abnormalities. This can be explained by the shared fetal circulation of monozygotic twins in utero. The group went on to corroborate their findings with a larger cohort of 58 SRS patients where 64% of the patients showed hypomethylation of the ICR1 region, a finding not seen in any of 68 SGA patients.

The resulting growth restriction could be caused by over expression of either CDKNIC, a known negative regulator of cell proliferation and growth, or PHLDA2, which has been found to be upregulated in IUGR babies.

Eggermann et al. had previously described two SRS patients with maternal duplication of 11p15 material and, together with the observation of epigenetic defects in this region of BWS patients, prompted Gicquel et al. to examine their SRS patients to see whether epigenetic defects may also explain the opposite growth phenotype. No methylation changes were seen in the centromeric ICR2 KCNQ1OT1 gene for any of their patients, but partial hypomethylation was observed in the H19 gene in

![Figure 1](http://jmg.bmj.com/)

**Figure 1** Role of methylation in regulation of IGF2 and H19 expression.

In normal individuals, CTCF binds to the unmethylated maternal allele ICR1 which prevents IGF2 promoters interacting with enhancers and silences IGF2. On the paternal allele, CTCF is inhibited from binding to the methylated ICR1, permitting IGF2 promoter association with enhancers which results in paternal allele expression only. In Beckwith–Wiedemann syndrome (BWS) patients, hypomethylation of the ICR1 permits IGF2 promoter association with enhancers from both maternal and paternal alleles resulting in over-expression of IGF2. In Silver–Russell syndrome (SRS) patients, hypomethylation of the ICR1 prevents IGF2 promoter interactions with enhancers resulting in no expression of IGF2 from either allele.

- **Table 1** Methylation analysis of ICR1 and ICR2 in growth affected patients

<table>
<thead>
<tr>
<th>No. patients studied</th>
<th>No. patients affected</th>
<th>ICR1 methylation</th>
<th>ICR2 methylation</th>
<th>Notes of interest</th>
<th>Refs</th>
</tr>
</thead>
<tbody>
<tr>
<td>9 SRS</td>
<td>5/9 SRS</td>
<td>Hypomethylation of H19 DMD/ICR1</td>
<td>Normal</td>
<td>Unaffected monozygotic twin; no hypomethylation in fibroblasts</td>
<td>53</td>
</tr>
<tr>
<td>3 Con 1 BWS</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3 SRS 7 IUGR 2 Con 2 BWS</td>
<td>2/3 SRS 7/7 IUGR</td>
<td>Examined H19 promoter only. Complete hypomethylation of two SRS patients</td>
<td>Normal</td>
<td>One additional SRS patient had no methylation aberration</td>
<td>69</td>
</tr>
<tr>
<td>51 SRS 69 SGA (Non-SRS)</td>
<td>16/51 SRS 37/58 SRS</td>
<td>Hypomethylation H19 DMD/ICR1: MI 20–37%</td>
<td>Normal</td>
<td>80% of patients with hypomethylation defects had body asymmetry</td>
<td>70</td>
</tr>
<tr>
<td>58 SRS</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 SRS 1 SRS</td>
<td>1 SRS</td>
<td>No methylation defects noted</td>
<td>Maternal ICR2 duplication</td>
<td>Normal IGFII serum in 82 patients mUPD 7 excluded; paternally expressed KCNQ1OT1 including maternally expressed KCNQ1, KCNQ4, SLC22A8, PHLDA2</td>
<td>71</td>
</tr>
</tbody>
</table>

BWS, Beckwith–Wiedemann syndrome; Con, controls; IUGR, intrauterine growth restriction; MI, methylation index; SGA, small for gestational age; SRS, Silver–Russell syndrome.
types in BWS and SRS patients on chromosome 11 and have shown that the errors are of opposite parental origin including mutations, duplications and methylation deregulation.

There have been several recent reports of children born with SRS who had been conceived by assisted reproductive technology (ART). Two of these cases were found to have methylation defects. The ART embryo culture medium is a cocktail of growth factors which could conceivably have an effect on methylation of the embryo and hence its growth potential. Certainly studies on in vitro culture conditions on mice pre-implantation embryos have demonstrated long term physiological, neurodevelopmental and behavioural effects. Some of these changes may be due to the observed changes in mRNA expression of imprinted genes which are particularly vulnerable during the blastocyst stage. Although ART is generally considered safe, babies with low birth weight are over represented in this population of births, as well as children with BWS and Angelman syndrome (AS) caused by imprinting defects, and it is therefore of clinical significance to determine the optimal culture conditions for future ART progeny. This may be especially important when culture media contains methyl donors which may influence methylation and imprinting. There is also the consideration that ART is performed on patients with fertility problems which could carry a higher risk of passing on aberrantly imprinted genes to their offspring.

Kobayashi recently described abnormal paternal methylation imprints (14.5%) and abnormal maternal imprints (20.6%) in oligospermic men.

**Other chromosomes**

Apart from chromosomes 7 and 11, six other chromosomes have been associated with the SRS phenotype (table 2). In some instances the patients described have had additional or atypical features.

**Table 2: Other chromosomes (Chr) considered as candidate loci for Silver–Russell syndrome (SRS)**

<table>
<thead>
<tr>
<th>Chr</th>
<th>No. reports</th>
<th>No. patients</th>
<th>Genotype</th>
<th>Atypical features</th>
<th>Refs</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>2</td>
<td>1</td>
<td>Trisomy 1q32.1-42.1</td>
<td>None</td>
<td>79</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Dup 1q42-qter</td>
<td>None</td>
<td>80</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>1</td>
<td>1</td>
<td>Del 8q11-q12</td>
<td>Microcephaly, shallow supraorbital ridges, hypoplastic alae nasi</td>
<td>81</td>
<td></td>
</tr>
<tr>
<td>15</td>
<td>3</td>
<td>1</td>
<td>r15</td>
<td>Mental retardation, cardiac defects</td>
<td>82</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>r15 (p12q26.3)</td>
<td>Mental retardation, microcephaly</td>
<td>83</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Trisomy7q34-qter, monosomy 15q26.3-qter</td>
<td>Bilateral congenital glaucoma</td>
<td>84</td>
<td></td>
</tr>
<tr>
<td>17</td>
<td>3</td>
<td>1</td>
<td>t(17;20)(q25;q13)</td>
<td>None</td>
<td>88</td>
<td>A putative candidate gene in this region is choricin somatomammotropin 1 (CSH1) which was deleted in two SRS patients. 76–82 Growth factor receptor binding protein (GRB) 2 (17q24-25) and 7 (17q21-22) are also localised to this breakpoint. However, Hitchins et al investigated CSH1, GRB2 and GRB7 in 48 non-UPD 7 SRS patients with no abnormal findings</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>t(1;17)(q31;q25)</td>
<td>None</td>
<td>89</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Hypomethylation at 11p15 plus deletion of CSH1</td>
<td>None</td>
<td>90</td>
<td></td>
</tr>
<tr>
<td>18</td>
<td>3</td>
<td>1</td>
<td>Trisomy 18</td>
<td>Many extra features of trisomy 18 syndrome, mental retardation</td>
<td>94</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Trisomy 18</td>
<td>None</td>
<td>95</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>del 18p</td>
<td></td>
<td>96</td>
<td></td>
</tr>
<tr>
<td>X</td>
<td>2</td>
<td>1</td>
<td>45X right side</td>
<td>None</td>
<td>97</td>
<td>Partington et al described a family with short stature and abnormal pigmentation with symptoms more pronounced in males suggesting X-linked inheritance. Younger male siblings showed classical SRS features</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>45X/45XY left side</td>
<td></td>
<td>98</td>
<td></td>
</tr>
</tbody>
</table>

**CONCLUSION**

It is now well established that 10% of SRS patients have mUPD for chromosome 7 and up to 50% have methylation defects in the imprinted domain on chromosome 11p15. This still leaves at least 40% of SRS patients with an unknown genetic aetiology. This may be explained by patient ascertainment, where many of the diagnostic criteria are shared with other syndromes. Clinicians generally agree that there are both SRS patients and SRS-like patients with a grey area between. It may be that the diagnostic criteria need to be tightened to ensure that the classical cases are identified and separated from the non-classical cases. It will be useful to make a detailed comparison of the clinical features of all SRS patients in order to identify whether the patients can be further categorised. For example, this might include those with and without limb asymmetry and to correlate genetic aetiologies with specific symptoms. For such a genetically diverse syndrome it is still conceivable that multiple genetic defects per patient could also contribute to the final phenotype.

Epigenetics is an exciting field and one which will undoubtedly prove vital in explaining many more syndromes and diseases. Various forms of cancer have for some time been linked to epigenetics, but conditions affecting neurology, behaviour and psychopathology such as schizophrenia and Alzheimer’s disease have recently also been suggested to have epigenetic components. Furthermore, the finding that diet and environment can influence epigenetic outcome, both in utero and throughout life, makes for a challenging but promising field of study. It is clear that epigenetics plays an important role in fetal growth and development but the full understanding of the mechanisms and their effect on gene function is yet to be elucidated, particularly for the benefit of clinical application. Human epigenetic models such as SRS will continue to aid us in this pursuit.


