Renal transplantation in infants

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Abstract Renal transplantation (RTx) has become an accepted mode of therapy in infants with severe renal failure. The major indications are structural abnormalities of the urinary tract, congenital nephrotic syndrome, polycystic diseases, and neonatal kidney injury. Assessment of these infants needs expertise and time as well as active treatment before RTx to ensure optimal growth and development, and to avoid complications that could lead to permanent neurological defects. RTx can be performed already in infants weighing around 5 kg, but most operations occur in infants with a weight of 10 kg or more. Perioperative management focuses on adequate perfusion of the allograft and avoidance of thrombotic and other surgical complications. Important long-term issues include rejections, infections, graft function, growth, bone health, metabolic problems, neurocognitive development, adherence to medication, pubertal maturation, and quality of life. The overall outcome of infant RTx has dramatically improved, with long-term patient and graft survivals of over 90 and 80%, respectively.

Keywords Infant · Kidney transplantation · Growth · Infections · Neurocognitive development

Introduction

The first kidney transplantations (RTx) to infants (0–24-month-old children) were performed in the 1960s in Minneapolis [1, 2]. The long-term results were at first modest and skepticism was expressed on the rationality of treating small children with severe renal failure. In the 1980s, technical refinements in dialysis therapy and post-RTx immunosuppressive medication dramatically improved the overall results of renal replacement therapy (RRT) in children and adults, and also led to a gradual acceptance of these therapies for infants with end-stage renal disease (ESRD).

Infants still form a minor group in pediatric RTx programs. The incidence of children needing RRT before the age of 2 years is low (7–8 per million age-related population) [3]. In a survey from the UK and Ireland, infants accounted for only 2.8% (19/675) of pediatric patients with ESRD [4]. In the North American Pediatric Renal Trials and Collaborative Studies (NAPRTCS) registry data on 10,632 RTx recipients, children <24 months accounted for 5.3% of pediatric recipients [5]. In a recent report from Europe, Australia, New Zealand, and Japan, very few of the children who already needed RRT during the neonatal period were transplanted before the age of 12 months, and 80% of these children were still on dialysis at the age of 24 months [3]. There is, however, center-based variation and many large specialized centers perform RTx for younger infants.

Recipient characteristics

The major indications for infant RTx are congenital abnormalities of the urinary tract (CAKUT), congenital nephrotic syndrome (CNS), neonatal cortical necrosis due to thrombosis, and autosomal recessive polycystic kidney disease (ARPKD) [6]. A small proportion of patients have a diagnosis seen in
older children, such as hemolytic uremic syndrome and oxalosis. It is important to note that acute kidney injury in the neonatal period, particularly in premature infants, often recovers and rarely leads to ESRD and RTx in infancy. Similarly, most infants with CAKUT show slow progression and the patients manage into adolescence or adulthood without RRT [7]. The progression of renal failure in inherited CNS may also be slow and RTx can be performed at the age of 2–5 years.

Active treatment of infants before RTx is important to ensure optimal growth and development as well as to avoid complications (thromboses, septic infections) that could lead to permanent neurological defects. Nutritional therapy, adequate calcium and vitamin D supplementation, phosphate-binding medication, anemia and infection, as well as management of hypertension and proteinuria are required to prevent the progression of renal failure [8]. Peritoneal dialysis (PD) is mostly used for ESRD management before RTx [9] and hemodialysis (HD) is reserved for those with problems in PD. However, in patients with oxalosis, organic acidemias, extensive history of abdominal surgery, poor socioeconomic circumstances, and fear for familial non-adherence, HD is the preferred treatment modality. Home PD in an anuric infant is also demanding, and improvements in pediatric HD may increase its use in coming years.

Absolute contraindications for RTx are uncontrolled malignancy and infection, as is the case with older patients. Children with a history of Wilms tumor who traditionally have been transplanted 2 years after the end of cancer therapy (European guidelines), but recent data suggest that patients with a low-risk tumor who remain relapse-free for 6 months after treatment might be considered for transplantation [10]. Up to 40 % of infants with ESRD have CAKUT and the anomalies, which were previously regarded as a contraindication to RTx, may affect the long-term kidney graft function and survival. Preoperative assessment of the bladder function with urodynamic studies is crucial in CAKUT patients. No consensus exists whether augmentation of the bladder should be performed pre- or post-transplant in children needing it. In our unit, infants usually undergo ureterocystoplasty before transplantation, whereas intestinal cystoplasty and Mitrofanoff conduit are performed after RTx, if needed. According to recent reports, CAKUT patients manage quite well after RTx and augmentation does not seem to affect the overall graft or patient survivals [11–14].

More than one-third of pediatric patients with ESRD have co-morbidities, such as chromosomal abnormalities, syndromic diagnosis, cerebral palsy, heart disease, and developmental delay, which may severely worsen the life quality and overall prognosis after RTx [4]. Withholding treatment from these infants is a difficult ethical question with no simple answer [6]. In general, attitudes have shifted with time and more handicapped patients are currently accepted for RRT. An individualized decision is usually made by a multidisciplinary team from the transplant center before the commencement of chronic dialysis.

The practice in most transplant centers is to start chronic dialysis only in those infants who are regarded as candidates for transplantation. The crucial question is how and to what extent the child and the family benefit from the operation [15].

Donor characteristics

A kidney from either of the parents is often used in infant RTx. According to a NAPRTCS report from the US, a living donor (LD) kidney was used in about 50 % of all pediatric RTx and in 74 % of infant RTx [5]. In the Eurotransplant region, the percentage of LD in pediatric RTx was much less (20 %), but no exact figures were available for infant RTx [16]. The use of either of the parents as an LD brings several advantages: timing of the operation is feasible, RTx can be performed pre-emptively, immunological matching is at least satisfactory (one haplotype), the donor age is often low (<35 years), and the cold-ischemia time is very short. On the other hand, the use of a young parent for a kidney donation may raise ethical concerns on the safety of the operation and the long-term health of the donor. It has been shown that living kidney donors maintain long-term renal function and experience no increase in cardiovascular or all-cause mortality. However, in recent surveys, donors showed a slightly increased long-term risk for ESRD and cardiovascular complications [17, 18].

Most genetic kidney diseases manifesting soon after birth show autosomal recessive inheritance and either parent can be used as a donor also in these diseases, such as CNS and ARPKD. If the disorder is not recessively inherited, genetic consultation is important to prevent recurrences after RTx.

The allocation of deceased donor (DD) kidney grafts to children varies worldwide. In the US, the current allocation policy for children (Share35) emphasizes the importance of young donors (<35 years old) and shorter waiting times over HLA-matching [19]. This algorithm has shortened the waiting times but increased mismatches, so that 84 % of transplants to children are mismatched at 4–6 alleles (A,B,DR loci) [20]. A poor match in the first graft also impairs the chances for the second graft, which is important in infant RTx, as these patients inevitably need another graft later in life [21]. In most European centers HLA-A, B, and DR matching is still an important part of organ allocation [19, 22], which may be reflected in the better long-term survival rates in Europe [23]. The great majority of kidney grafts come from brain-death donors, and the experience of donation after cardiac death (DCD) is still limited in pediatric and especially infant RTxs [24].

Traditionally, kidneys from young pediatric donors (<5 years of age) have not been used in infant RTx, as the early reports showed a decreased graft survival rate caused by infections and technical problems [25–28]. However, more recent data indicate that a pediatric donor organ can adapt the glomerular filtration rate (GFR) with the recipient’s
growth and these grafts would provide even better long-term results [29, 30]. En bloc kidneys from infant donors have been used for adult recipients and more recently also for older pediatric recipients (>4 years of age), but, to our knowledge, not for infant recipients [31].

ABO-blood group compatibility between the donor and recipient has traditionally been required in RTx. However, ABO-incompatible (ABOi) renal transplantation has become more popular and hundreds of patients have been successfully transplanted across the ABO-barrier [32–34]. Infants have low levels of ABO-antibodies and are immunologically good candidates for ABOi-transplantation. This is evident in heart transplantations, where most ABOi recipients are infants [35]. Thus, it is to be expected that the use of ABOi donors will increase in infant RTx during the coming years.

Timing of transplantation

Timing of the infant RTx is an important issue. Early transplantation is favored by the fact that transplanted children show better growth, development, and quality of life compared to dialysis patients, and they avoid many complications, such as peritonitis and venous access problems. Both intra- and extraperitoneal placement of the graft can be used (see below) and young infants weighing <5 kg have been successfully transplanted [1, 36, 37]. Very early transplantation carries, however, added risks and needs multidisciplinary expertise. Adult-sized grafts have been successfully transplanted into infants weighing 7–10 kg, but a common practice is to perform the operation when a child has reached the weight of 10–20 kg when extraperitoneal placement of the kidney is feasible [36]. This approach was evident in two large international surveys of about 400 infants with ESRD [3, 38]. Besides weight, the child’s height, body shape, and size of blood vessels play a role in the planning of the operation. In patients suitable for RTx, long dialysis period does not bring additional value and an early operation is preferred, especially in large specialized transplant centers.

In CNS patients with severe proteinuria, bilateral nephrectomy can be performed at the age of 6–9 months followed by dialysis therapy for some months before an early RTx (weight around 10 kg). The other strategy is to perform a unilateral nephrectomy and wait for the development of ESRD. In this case, RTx can be postponed to the age of 2–5 years [39]. According to a recent survey of EDTA-registry data, both approaches seem to result in equally good long-term outcome (T. Hölttä, personal communication, March 20, 2015).

Surgery

In infant RTx, a kidney graft can be placed intra- or extraperitoneally (Fig. 1 and Table 1). In the former approach, a midline incision is made on the abdominal wall and the graft is placed into the right side of the peritoneal cavity [1, 2]. After mobilizing the right colon and terminal ileum, the donor renal vein and artery are anastomosed to the recipient vena cava and aorta, respectively, in an end-to-side technique. The donor’s ureter is implanted into the recipient’s bladder using either a Ledbetter–Politano procedure or one of its modifications. Temporal stenting may be used to reduce the risk of ureteral stenosis, especially when suboptimal arterial blood flow in the distal ureter is suspected. An adult-sized kidney can occupy the entire right side of the abdomen leading to bowel dysfunction and ileus, especially during the early post-operative phase. The lateral edge of the allograft is usually easily accessible to an ultrasound-guided renal biopsy [1, 2, 40].

When an extraperitoneal placement is used, an incision is made above the right groin (or left side in re-RTx). The peritoneum is mobilized from the anterolateral and posterior abdominal wall, exposing the posterior muscles and the great vessels. The kidney graft artery and vein are usually anastomosed with the common iliac artery and vein, respectively. The sites for anastomoses, however, depend on the relative sizes of the vessels of the graft and the recipient. The greater the size mismatch is, the more proximal recipient vessel anastomoses are used. Still, the available space for an adult-size kidney may present a problem, and to avoid pressure and circulatory problems, it is sometimes safer to close the fascia a few days after the primary operation [30, 41]. To ensure urine flow, a transvesical catheter is usually placed in the bladder. In infants with CAKUT, there are different intra- and extravesical techniques to make a uretero-bladder anastomosis; the choice depends on the anatomy and previous surgery [13].

Peri- and postoperative management

While intraoperative management of the recipient follows normal anesthetic practice, maintenance of sufficient perfusion of the transplanted kidney is crucial. Sufficient cardiac output and volume overload is needed to ensure adequate perfusion of the allograft. The cardiac output of infants must double in order to perfuse the adult kidney adequately [42]. It is desirable to maintain a central venous pressure above 10 cm H₂O prior to unclamping and the mean arterial pressure at more than 60 mmHg. Intravenous crystalloid or colloid solutions (Ringersteril, 0.9 % normal saline, 4 % albumin) and mannitol may be used to promote urine output according to the practice of the transplant center.

After the operation, attention to the intravascular volume and electrolyte and acid–base stability is essential to ensure good renal function. During the first days, polyuria is common and urine output can be replaced by 0.45 % (or 0.9 %) saline solution. Additional fluid infusions may be given if urine output drops. Intravenous furosemide boluses can be given if
dehydration is excluded. Weight measurements, blood pressure monitoring, and follow-up of fluid input and output are important for securing the safety of excessive fluid administration (2500 ml/m²) [42]. Infant recipients are often slightly oedematous (0.5–1.0 kg) and hypertensive during the first postoperative days. Systolic blood pressure values of 100–120 mmHg are, however, allowed in this early phase.

Monitoring of serum sodium, potassium, bicarbonate, calcium, phosphorous, and magnesium is essential. Due to variation in the urine output (heavy diuresis, delayed graft function) and use of diuretics and calcineurin inhibitors (CNI), any of these electrolytes may need supplementation. On the other hand, slight hyperkalemia (5–6 mmol/l) is very common during the first weeks but rarely needs therapy. Oral phosphate and magnesium supplementations may be needed for several months, and calcium-vitamin D supplementation can be given permanently.

Hypertension secondary to high fluid intake, corticosteroids, and CNI is common after RTx. Also, the preoperative blood pressure of both the donor and recipient have an impact on the post-transplantation blood pressure values. Calcium-channel blockers (amlodipine and nifedipine) are widely used in pediatric RTx patients. Antibiotic prophylaxis is given as long as the bladder catheter is kept in situ. Infants without urine output before the operation have a small shrunken urinary bladder, and the catheter is closed for increasing time periods before its final removal. The use of anticoagulation (enoxaparin, ASA) postoperatively is decided individually. Our practice is to start anticoagulation if poor blood flow in the ureter is suspected or a small pediatric graft is used. Prophylaxis against Pneumocystis jirovecii (co-trimazole) and cytomegalovirus (CMV) (ganciclovir/valganciclovir) are routinely used by most centers for 3–12 months after the operation. Infants are often CMV negative at the time of RTx and receive a CMV positive graft from an adult donor and, thus, have a clear risk for CMV infection.

Immunosuppressive medication protocols are center-specific. This is so for both perioperative induction therapy and maintenance medication. In the NAPRTCS report from 2010, no induction was used in 54 %, antithymocyte globulin (ATG) in 27 %, and basiliximab (anti-IL2-receptor antibody) in 10 % of the US centers [5]. Triple medication with CNI (tacrolimus or cyclosporine A (CsA)), antimetabolite (mycophenolate mofetil (MMF) or azathioprine) and glucocorticoid (prednisone or methylprednisolone) is typically used as an early maintenance medication. According to the NAPRTCS report, immunosuppression at 30 days after RTx included prednisone in 50 %, tacrolimus in 60 %, MMF in 60 %, and CsA and azathioprine in only a small percentage in the US centers [5].

In general, infants need the same medication as older pediatric RTx patients. In our center, the early immunosuppressive protocol is still CsA-based, as the dosing in the smallest recipients is easier and more accurate with CsA as compared to tacrolimus. Due to the faster metabolic capacity of smaller children, CsA is at first given to the infants three times a day [43]. Steroid-free protocols are successfully used in many transplant centers to avoid the side effects of glucocorticoids.
The incidence of thrombosis has been 2–30 % were reported, but recently the recipients is graft thrombosis. In the original reports on infant RTx, frequencies up to 30 % were reported, but recently the occurrence of thrombosis has been 2–10 % in infants. The smaller size of the blood vessels predisposes infants to thrombotic events, and investigation of the size of blood vessels in the recipient before RTx is recommended. Also, perfusion and reimplantation damage, long ischemia time, hypotension and hypoperfusion of the graft, as well as immune mechanisms have been associated with the development of thrombosis.

The literature on acute antibody-mediated rejection in pediatric and especially infant RTx is still scarce. It is to be expected that preformed HLA-antibodies are less common in infants than in older children. De novo donor-specific antibodies (DSA) can be detected also in pediatric patients and they have been associated with decreased long-term graft function. However, contradictory results on the importance of DSA have been published.

Acute rejection

Improvements in immunosuppressive medication have resulted in a remarkable decrease in the incidence of acute rejection (AR) in pediatric RTx patients. In the latest NAPRTCS-registry (years 2007–2010), the probability of AR during the first 12 months had decreased to 8.6 and 16.6 % in LD and DD transplantations, respectively. An interesting question is whether infants have a different risk for rejection as compared to older children. Previously, heightened immunoactivity was suggested. However, in the recent registry data, infants showed relative hazard rates of 0.51 and 0.89 for the first AR in LD and DD transplantations, respectively, indicating that the risk was moderately decreased. In our experience, infants are not very different from older children in developing AR. A renal biopsy is mandatory for a diagnosis of AR in infants, since a large adult-sized kidney can undergo substantial damage before a rise in serum creatinine occurs.

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Infections

Peri- or postoperative bacterial or yeast infections are not common in infant RTx recipients. Urinary tract infections (UTI) are, however, quite frequent and associated with vesicoureteral reflux to the graft and sometimes to native kidneys. In the study of Their et al., the frequency of UTI in RTx children <2 years and 2–5 years of age was 0.3 and 0.1 episodes/patient year, respectively, suggesting that infants are not especially prone to UTI. Increased occurrence of UTI has been reported in RTx patients with CAKUT and bladder dysfunction. In recent reports, the UTI frequency in patients with diverted or augmented bladder was, however, not exceptionally high. Management of UTI is important, as repeat episodes may hasten deterioration of graft function and with some infants it is wise to use antibiotic prophylaxis.

Viral respiratory tract infections (URI) are by far the most frequent infectious complication of RTx infants. In our experience, complicated respiratory infections among RTx recipients are, however, rare, and even small children recover from URI quite normally; hospitalization of a patient due to URI is rarely needed. As is the case in the general population, gastrointestinal infections are not rare among small transplant children. Diarrhea and vomiting may require intravenous fluid therapy and medication as well as intensified drug therapy.
monitoring. Even a moderate dehydration often leads to temporally increased creatinine levels in RTx infants.

CMV, Epstein–Barr virus (EBV), and Polyomaviruses (PV) (nephritis caused by BK virus and less often JC virus) are a major concern in pediatric RTx patients of all ages. Since most infants are seronegative for these viruses at the time of transplantation, primary infection after RTx is common. Monitoring of these viruses by a PCR method from blood or plasma samples is highly recommended [57]. CMV viremia is rare in patients receiving valganciclovir prophylaxis, but after stopping the medication, subclinical or clinical viremia is quite common.

Management of PV and EBV viremias is more complicated, as there is no effective therapy for either virus, with the exception of reduction of the immunosuppressive medication. In the case of PV infection, nephritis caused by this virus can be verified by kidney biopsy, which should be taken when significant viremia (>10,000 copies/ml) or signs of renal graft dysfunction are noticed. Constant EBV viremia may lead to the development of post-transplant lymphoproliferative disease (PTLD) and increasing EBV DNA levels usually require reduction of immunosuppressive medication, which increases the risk of rejection [58]. In a recent survey, the relative hazard for PTLD was 5.3-fold higher in children aged <6 years of age versus those >12 years, and EBV seronegative subjects had a 4.7-fold higher risk compared to EBV-positive subjects [58]. In our center, PTLD has been diagnosed in seven of the 252 RTx patients (3.7 %) and four of these had received their kidney graft as infants. Excellent articles on the management of CMV, EBV, and PV infections have recently been published [57–61].

Long-term problems

While the short-term results of infant RTx are nowadays excellent, the major issue is how these children manage in the long run (Table 2). Patients who received RTx at young age need follow-up and therapy for several decades, not several years. Important issues are long-term graft function, growth, pubertal maturation and fertility, bone health, metabolic problems, neurocognitive development, adherence to medication, and quality of life.

Graft function

The most important cause of late graft failure is chronic renal allograft injury that accounts for one-third of the graft losses in pediatric RTx patients [5, 62]. The term includes chronic active T-cell-mediated and antibody-mediated rejections, arteriopathy and interstitial fibrosis/tubular atrophy with no specific etiology [63, 64]. According to our experience, chronic allograft injury is detected to a similar extent in the allografts of patients who received their transplant in infancy or later in childhood [65]. This is reflected in the similar pattern of GFR deterioration with time, as shown in Fig. 2. The mean annual decline of measured GFR is 2.2 ml/min/1.73 m², indicating that infant RTx recipients need a new graft as young adults.

Growth

Most infants are growth-deficient at the time of RTx. The NAPRTCS registry data showed that the youngest recipients (1–2 years) had a negative height Z score (hSDS) of −2.2 at the time of transplantation, but exhibited substantial catch-up growth (up to −1.4 SD) for the initial 3–4 years after transplantation before plateauing and subsequently exhibiting a decline in the height score (−1.7 SD at 6 years) [66]. In the study by Qvist et al., children <2 years of age had a mean hSDS of 1.1 both at RTx and 7 years after the operation [67]. Growth after RTx is satisfactory, but the final height attained by most recipients is not their calculated target height [68, 69]. The results

![Fig. 2 Measured annual glomerular filtration rate (GFR) in 61 patients transplanted before the age of 24 months and in 127 patients transplanted after the age of 24 months (Children’s Hospital, Helsinki)](image)
emphasize the importance of the preoperative management of infants with ESRD or nephrosis [70]. After RTx, allograft function and steroid exposure have an impact on growth, and in those with poor growth, steroid dosing should be minimized. Especially in adolescence, short stature can have major consequences on quality of life and self-esteem.

For the most stunted patients, recombinant growth hormone (GH) therapy has been used for over 20 years, and the therapy has been effective in inducing substantial catch-up growth in most patients. There have, however, been some concerns that the therapy might induce rejections or PTLD. A recent meta-analysis confirmed that GH therapy promotes growth velocity in RTx children [71]. In this analysis, the risk for rejection was slightly increased (risk ratio 1.56) in patients receiving GH. On the other hand, a report from Australia and New Zealand did not find a relationship between GH use and PTLD [72]. Monitoring of growth for at least 1 year post-transplantation is recommended before starting (or restarting) GH therapy. One practical problem with GH therapy is that it is expensive, and must be continued for years, as the positive effect stops when the therapy is discontinued.

**Pubertal development**

Besides growth, pubertal developmental and fertility are important long-term issues in adolescents and adults who were transplanted as infants. Delayed puberty has been reported in several studies on pediatric RTx patients [73, 74]. In a recent report by Tainio et al., children who underwent RTx at a young age (most patients <2 years at RTx) had quite normal pubertal development and reached it earlier than those transplanted at later age (12.3 vs. 13.4 years) [69]. Twenty percent of boys and none of the girls had a delayed onset of puberty. The bone age was delayed in practically all and final height was achieved at age 18.1 and 16.0 years in boys and girls, respectively, which provides growth potential for a longer time. In a subsequent study, the reproductive endocrine function of adult men who received RTx as small children was, however, impaired. Despite quite normal sex hormone levels, only one-fifth of these young men had normal sperm counts [75]. Studies on the fertility of young women transplanted in early childhood are still required.

**Bone health**

Mineral and bone disorder in RTx children may result in decreased bone mineral density (BMD), fractures, bone pain, and growth failure [76, 77]. In a cross-sectional study by Valta et al., vertebral fractures were observed in 8% of RTx children (median age 12 years) and the majority of them were asymptomatic. The height-adjusted BMD in lumbar spine was satisfactory and similar in patients transplanted before the age of 2 years as compared to those transplanted at a later age (−0.5 SD vs. −0.4 SD) [78]. Female sex and age >15 years as well as high PTH levels were significant predictors of low BMD. As the basis for lifelong bone health is established in childhood and adolescence, follow-up measurement of bone mineralization by DXA and spinal imaging of vertebral fractures is warranted [79]. Therapeutic efforts to reduce MBD include vitamin D, calcium, and sometimes phosphate supplementation.

**Metabolic risk factors**

In children transplanted at a very young age, avoidance of metabolic risk factors such as obesity, hypertension, dyslipidemia, and impaired glucose metabolism, is important in two respects. First, these factors may adversely affect the graft function and, secondly, they may lead to early cardiovascular problems that can impair later therapies (retransplantation in adulthood). The data on metabolic risk factors among pediatric RTx patients are still limited. In a cohort of patients mostly transplanted before the age of 2 years, metabolic syndrome, overweight, hypertension, and type 2 diabetes were observed in 14–19%, 20–23%, 61–87%, and 3–5% of patients at 1.5–5 years after RTx, respectively [45]. Higher incidences of these risk factors, however, have been reported in pediatric RTx patients [80–83]. Thus, the follow-up of blood pressure is crucial and antihypertensive medication is required in the majority of the patients. In addition, significant proteinuria is associated with worse outcome and should be taken into account in the medication [84].

**Neurocognitive development**

Children with renal failure from infancy would be expected to have a less favorable neurodevelopmental prognosis. This is especially so in patients with neurological comorbidities and those who suffered from thrombotic events before RTx. However, in many children, the neuromotor development after RTx is satisfactory. In a recent report, patients (mean age, 11 years) diagnosed with ESRD as infants had intellectual and metacognitive functioning significantly lower than sibling controls with the mean Full Scale IQ scores of 78 and 94, respectively [85]. In a report of 33 school-aged children transplanted before the age of 5 years, younger age at RTx was associated with higher scores on several parameters. The mean intelligent quotient (IQ) was 87, and 6–24% showed impairment in neuropsychological tests. About 80% of the children attended normal school and 76% had normal motor performance [86]. In another study, 50 children transplanted at an early age were assessed at a mean age of 11.1 years. The RTx group scored generally lower than the control group on neuropsychological assessment. The difference was evident in both the verbal and visuospatial domains and verbal working memory. A better cognitive outcome was associated with the...
absence of neurological co-morbidity, younger age, shorter disease duration, and sustained kidney function [87].

Life quality

The ultimate goal of RTx is to provide a good overall well-being of the patients. In general, the psychosocial outcome after RTx children is satisfactory, so that most patients and their parents have a good life according to themselves [88]. In a study of psychosocial adjustment in school-aged children, most of whom had received an RTx during infancy, the health-related measure was comparable to normal school children [89]. The reported results have, however, been variable, and in a quite recent study, RTx children (aged 3–19 years) showed higher levels of mental health problems and lower quality of life as compared to controls [90]. Quality of life has also been studied in young adults who received RTxs in childhood. Again, the results are varied, but most patients are quite satisfied with their life [91–93]. Adolescents who received RTxs in infancy, and do not remember the “hard times” before and after RTx, need thorough surveillance for adherence to medication and help in transition to an adult unit.

Comorbidities significantly affect quality of life and many disabilities originate from the time before RTx [94]. Although the quality of life is not necessarily correlated to the degree of physical disability, much effort should be made to diminish the complications, which have an impact for later adjustment. Continuous multidisciplinary support, follow-up, and education are needed to cope with this problem.

Key summary points

– In infants, congenital anomalies of the kidneys and urinary tract (CAKUT) and congenital nephrotic syndrome are the most common causes of renal transplantation. Active treatment of an infant with renal failure is important to ensure optimal postoperative outcome.
– Renal transplantation with intraperitoneal engraftment can be performed on an infant weighing 6–10 kg. If the graft is placed extraperitoneally, transplantation is usually performed when the child has reached the weight of 10–15 kg. Intra- and perioperative management of a recipient is focused on maintenance of sufficient perfusion of the transplanted kidney and avoidance of thrombosis, rejections, and infections.
– Infants need life-long immunosuppressive medication, including calcineurin inhibitor, antimetabolite, and corticosteroid. Management of the many side effects of these drugs is important to ensure good growth and development.

Conclusions

The overall outcome of infant RTx has dramatically improved such that several registry data and single-center reports show 10-year patient and graft survivals of over 90 and 80 %, respectively [95]. Short-term problems after transplantation are more common in infants than in older children [5], but the long-term outcome figures in infants are better, with a relative hazard for graft failure being about 0.2–0.6 in infants as compared to adolescents [96]. Infants receiving living donor grafts have estimated graft half-life of almost 30 years [96]. Careful follow-up of subjects who received kidney grafts as small children is needed to ensure optimal growth and development, as well as good quality of life in adulthood when the second kidney transplantation is required.

Multiple-choice questions (answers are supplied following the reference list)

1) Infants form a special group of pediatric renal transplant recipients. Of all pediatric recipients they account for:
   a. <10 %
   b. 10–19 %
   c. 20–29 %
   d. 30–39 %
   e. >40 %

2) Acute rejections are nowadays diagnosed less often than before. Their frequency in infant renal transplantation is:
   a. 40–60 %
   b. 30–39 %
   c. 20–29 %
   d. 5–19 %
   e. <5 %

3) Infant kidney transplant recipients are often seronegative for Epstein–Barr virus (EBV) at the time of transplantation. The risk for post-transplant lymphoproliferative disorder (PTLD) in seronegative recipients, as compared to seropositive subjects, is:
   a. 2-fold
   b. 5-fold
   c. 8-fold
   d. 10-fold
   e. 20-fold

4) In infant renal transplantation the major cause of early graft loss is:
   a. acute rejection
   b. urinary tract infection
   c. ureteral stenosis
d. lymphocele
e. vascular thrombosis

5) The overall outcome of renal transplantation in infants is nowadays good. Graft survival ten years after the operation is:
   a. 50 %
   b. 60 %
   c. 70 %
   d. 80 %
   e. 99%

Conflict of interest  The authors declare that they have no conflict of interest.

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