Fetoscopic laser photocoagulation in twin-to-twin transfusion syndrome: experience from a single institution

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INTRODUCTION

Twin-to-twin transfusion syndrome (TTTS) is the most common serious complication of monochorionic (MC) twin pregnancies, with perinatal mortality rates of up to 90% if untreated. This study aimed to review the perinatal and perioperative outcomes of MC twin pregnancies treated for TTTS by fetoscopic laser photocoagulation (FLP) since its introduction at KK Women’s and Children’s Hospital (KKH), Singapore, in 2011.

METHODS

This was a retrospective review of five consecutive patients who underwent FLP of placental anastomoses for TTTS at KKH from June 2011 to March 2014. FLP was offered to patients who were diagnosed with TTTS of at least Quintero Stage II before 26 weeks of gestation. The main outcome measures were perioperative complications and perinatal survival rates.

RESULTS

Five sets of MC twin pregnancies underwent FLP during the study period – three pregnancies were diagnosed with Stage III TTTS and two pregnancies with Stage II TTTS. Median gestational ages at initial presentation, laser photocoagulation and delivery were 19 (range 17–20) weeks, 20 (range 19–23) weeks and 29 (range 28–34) weeks, respectively. One patient had bleeding into the amniotic cavity intraprocedurally. Overall, the perinatal survival rate, double-infant survival rate and survival rate for at least one twin were 60% (6/10 fetuses), 40% (2/5 twins) and 80% (4/5 twins), respectively.

CONCLUSION

FLP is a feasible treatment for TTTS, with minimal maternal complications. Perinatal survival rates of this patient group that was managed at our centre were comparable to those of international centres.

Keywords: fetofetal transfusion, fetoscopic laser photocoagulation, monochorionic
for evidence of resolution, persistence or recurrence of TTTS. Parameters measured included deepest vertical pool, presence or absence of fetal bladder, standard deviation score of abdominal circumference, and Doppler studies of the UA, DV and middle cerebral artery (MCA). Operative details, perioperative and postoperative complications, and pregnancy outcomes were reviewed.

The overall survival rate was calculated by taking the number of surviving fetuses divided by the total number (n = 10) of fetuses. The double-infant survival rate was the proportion of pregnancies in which both fetuses survived. The one-twin survival rate was the proportion of pregnancies in which one or both fetuses survived.

All procedures were performed in the Antenatal Diagnostic Centre of KKH. All the patients had local anaesthesia for the procedure. Antibiotic prophylaxis in the form of intravenous cefazolin 1 g was given to all patients 30 minutes before the procedure. An ultrasonography examination was performed prior to the procedure to confirm the viability of the twin pregnancy, determine the location of the placenta and cord insertions of both twins, and to plan for the site of entry of the fetoscope. Instruments used included a 2-mm fetoscope (HOPKINS® II Straight Forward Telescope 0°; Karl Storz, Tuttlingen, Germany), a diode laser with a 600-μm laser fibre (Dornier Medilas D MultiBeam; Dornier MedTech, Wessling, Germany), vascular access trocar and cannula (Terumo Radifocus 11F; Terumo Corporation, Tokyo, Japan). Under ultrasonographic guidance, the trocar and cannula were inserted into the amniotic cavity of the recipient twin using the Seldinger method after a small abdominal incision was made. The site of entry was dependent on the placental location as well as the placental cord insertions of both twins. Once inside the amniotic cavity, the amniotic fluid in the recipient sac to that of normal liquor volume. Fetal viability was confirmed with ultrasonography at the end of the procedure, and Doppler studies of the UA, DV and MCA of both twins were routinely performed. The patient was then admitted for an overnight stay and started on oral nifedipine 10 mg administered four times a day.

### RESULTS

A total of five patients who underwent FLP from June 2011 to March 2014 were reviewed in this study. Four of the five patients were referred to our institution after 19 weeks of gestation. Of the four referred patients, three were booked early in private local hospitals, while one was booked early in an overseas hospital. The fifth patient was booked early in our institution.

### Patient 1

A 34-year-old woman (gravida 4, para 0) with a history of three previous miscarriages was booked early with a private obstetrician initially. She was referred to our institution at 19 weeks of gestation for possible TTTS and was later diagnosed with Stage III TTTS at our centre. FLP was performed at 19 + 2 weeks of gestation, according to our local protocol. TTTS resolved after FLP and both twins continued to grow till 30 + 2 weeks. The ex-donor twin

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Patient 1</th>
<th>Patient 2</th>
<th>Patient 3</th>
<th>Patient 4</th>
<th>Patient 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maternal age (yr)</td>
<td>34</td>
<td>35</td>
<td>32</td>
<td>41</td>
<td>28</td>
</tr>
<tr>
<td>Gestation at diagnosis (wk)</td>
<td>19</td>
<td>20</td>
<td>17</td>
<td>20</td>
<td>19 + 2</td>
</tr>
<tr>
<td>Quintero stage at diagnosis</td>
<td>III</td>
<td>III</td>
<td>III</td>
<td>II</td>
<td>II</td>
</tr>
<tr>
<td>Gestation at FLP (wk)</td>
<td>19 + 2</td>
<td>23 + 2</td>
<td>21</td>
<td>20 + 3</td>
<td>19 + 3</td>
</tr>
<tr>
<td>Gestation at delivery (wk)</td>
<td>30 + 6</td>
<td>29</td>
<td>27 + 4</td>
<td>28 + 5</td>
<td>34</td>
</tr>
<tr>
<td>Mode of delivery</td>
<td>LSCS</td>
<td>LSCS</td>
<td>LSCS</td>
<td>LSCS</td>
<td>LSCS</td>
</tr>
<tr>
<td>Procedure-delivery interval (wk)*</td>
<td>8 (6–15)</td>
<td>8 (6–15)</td>
<td>8 (6–15)</td>
<td>8 (6–15)</td>
<td>8 (6–15)</td>
</tr>
</tbody>
</table>

FLP: fetoscopic laser photocoagulation; IUD: intrauterine death; LSCS: lower-segment Caesarean section; UD: vaginal delivery

<table>
<thead>
<tr>
<th>Perinatal outcome</th>
<th>No. (%)</th>
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<tbody>
<tr>
<td>Overall survivor (per fetus) (n = 10)</td>
<td>6 (60)</td>
</tr>
<tr>
<td>Double survivor (per pregnancy) (n = 5)</td>
<td>4 (80)</td>
</tr>
<tr>
<td>At least one survivor (per pregnancy) (n = 5)</td>
<td>4 (80)</td>
</tr>
<tr>
<td>Gestational age at delivery (wk)*</td>
<td>29 (28–34)</td>
</tr>
<tr>
<td>Procedure-delivery interval (wk)*</td>
<td>8 (6–15)</td>
</tr>
</tbody>
</table>

*Data presented as median (range).
suffered an unexpected intrauterine fetal death at 30 + 2 weeks (Day 77 post-FLP). Serial measurements of MCA-peak systolic velocity (PSV) were performed daily to monitor the status of the ex-recipient twin; the measurements were noted to be on a consistently increasing trend, from 1.11 multiples of median (MoM) to 1.20 MoM, and finally at 1.29 MoM over the three days following the demise of the ex-donor twin. Decision for delivery via elective Caesarean section at 30 + 6 weeks was made in view of the rising MCA-PSV in the ex-recipient twin, which indicated possible onset of fetal anaemia. Birth weight of the surviving twin was 1,505 g and the Apgar scores were the same at one and five minutes (i.e. 9). Haemoglobin level of the neonate at birth was normal, at 18.6 g/dL. The baby stayed in the neonatal intensive care unit for 13 days and the special care nursery for 28 days, and was discharged from hospital on Day 41 of life.

Patient 2
A 35-year-old woman (gravida 1, para 0) with a spontaneous monochorionic-diamniotic (MCDA) twin pregnancy was booked early at an overseas hospital initially. She was diagnosed with TTTS at 20 weeks of gestation. The patient decided to seek treatment at our institution and was booked at 23 weeks. She was diagnosed with Stage III TTTS at our centre and FLP was performed at 23 + 2 weeks. Laser ablation was performed on the identified anastomotic vessels. The procedure was complicated by haematoma formation at uterine-placenta wall insertion intraoperatively, and the liquor became bloodstained as a result. Laser fetoscopy was thereafter terminated and amnioreduction was performed, with 2,000 mL of bloodstained liquor reduced in total. The patient was admitted to the labour ward for monitoring. Repeat ultrasonography the following day noted cessation of fetal cardiac activity of the donor twin. Dopplers changes in the recipient resolved and the ex-recipient twin continued to grow. However, the ex-recipient twin died on Day 39 post-FLP at 28 + 6 weeks of gestation.

Patient 3
A 32-year-old woman (gravida 2, para 1) with a history of one previous lower-segment Caesarean section was booked early with a private obstetrician initially. She was diagnosed with TTTS at 17 weeks of gestation and referred to our centre at 20 + 1 weeks, where Stage III TTTS was diagnosed and FLP was performed at 21 weeks. TTTS resolved, but the ex-donor twin appeared to have Type 3 selective intrauterine growth restriction as defined by Gratacós et al.,(7-9) with the estimated fetal weight of the small fetus falling below the tenth percentile, and the presence of intermittent absent and reversed end-diastolic flow on UA Doppler ultrasonography. She was monitored closely in view of the abnormal Doppler changes. At 27 + 2 weeks of gestation, the ex-donor twin had reversed a-wave noted on DV Doppler ultrasonography and hence a decision was made for delivery. Intramuscular dexamethasone was administered and the patient delivered via elective Caesarean section at 27 + 4 weeks of gestation. Birth weights of the ex-recipient and ex-donor twins were 1,224 g and 773 g, respectively. The Apgar scores at one and five minutes for the ex-recipient and ex-donor twins were 9 and 9, and 5 and 8, respectively. The ex-recipient twin was discharged on Day 76 of life. However, the ex-donor twin died on Day 17 of life due to sepsis and prematurity.

<table>
<thead>
<tr>
<th>Characteristic</th>
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<th>Patient 3</th>
<th>Patient 4</th>
<th>Patient 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth weight (g) (donor/recipient)</td>
<td>1,505</td>
<td>398/760</td>
<td>773/1,224</td>
<td>780/–</td>
<td>1,200/2,000</td>
</tr>
<tr>
<td>Apgar score (donor/recipient)</td>
<td>9/9</td>
<td>–/9</td>
<td>5/9</td>
<td>8/–</td>
<td>NA</td>
</tr>
<tr>
<td>NICU admission</td>
<td>One twin</td>
<td>–/9</td>
<td>Both twins</td>
<td>One twin</td>
<td>NA</td>
</tr>
<tr>
<td>Survival up to 28 days</td>
<td>One twin</td>
<td>–/9</td>
<td>One twin</td>
<td>One twin</td>
<td>Both twins</td>
</tr>
<tr>
<td>NICU stay (day) (donor/recipient)</td>
<td>–/13</td>
<td>–/76</td>
<td>17/–</td>
<td>–/76</td>
<td>NA</td>
</tr>
<tr>
<td>LOS (day) (donor/recipient)</td>
<td>–/41</td>
<td>–/76</td>
<td>175/–</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td>Perinatal morbidity</td>
<td>Recipient: hypoglycaemia, NNJ, mild pulmonary stenosis, late-onset Group B streptococcal sepsis</td>
<td>Donor: hypoglycaemia, NNJ, RDS, PPHN, PDA, VSD, IH, ARDS (demised after 17 days of life)</td>
<td>Recipient: hypoglycaemia, RDS, PDA, IVH, ROP, NNJ</td>
<td>Donor: hyaline membrane disease, PDA, sepsis, NNJ, ileal stenosis with intestinal obstruction necessitating ileostomy, infantile-type PCKD</td>
<td></td>
</tr>
</tbody>
</table>

ARDS: acute respiratory distress syndrome; IUD: intrauterine death; IVH: intraventricular haemorrhage; LOS: length of hospital stay; NA: not available; NICU: neonatal intensive care unit; NNJ: neonatal jaundice; PCKD: polycystic kidney disease; PDA: patent ductus arteriosus; PPHN: persistent pulmonary hypertension of the newborn; RDS: respiratory distress syndrome; ROP: retinopathy of prematurity; VSD: ventricular septal defect
Patient 4
A 41-year-old woman (gravida 6, para 2) with a history of three previous miscarriages and two previous normal vaginal deliveries was booked early with a private obstetrician. This was a spontaneous MCDA twin pregnancy. She was diagnosed with TTTS at 20 weeks of gestation and referred to our centre the day after diagnosis. Stage II TTTS was diagnosed and FLP was performed at 20 + 3 weeks of gestation. However, TTTS persisted despite the performance of FLP; the donor twin continued to have an empty bladder on follow-up visits. The mother then underwent amnioreduction at 23 + 1 weeks of gestation. However, as there was no improvement post-amnioreduction, repeat FLP was performed at 24 + 3 weeks of gestation. Repeat FLP is one of the approaches taken for the management of recurrent TTTS (other options include amnioreduction, expectant management or cord occlusion), with the internationally reported rate of a repeat procedure being 29.6% according to a systematic review in 2012.106 However, the ex-recipient twin died two days after repeat FLP at 24 + 5 weeks (Day 30 post-first FLP). The ex-donor twin survived and its amniotic fluid volume improved, with visualisation of the bladder. The patient subsequently went into spontaneous preterm labour at 28 + 5 weeks of gestation and delivered a baby of birth weight 780 g and Apgar scores of 8 at one minute and 8 at five minutes. The baby had multiple medical problems but was discharged on Day 175 of life.

Patient 5
A 28-year-old woman (gravida 1, para 0) had booked early at our institution, at 12 weeks of gestation with a spontaneous MCDA twin pregnancy. She was diagnosed with Stage II TTTS at 19 + 2 weeks of gestation and underwent FLP the following day at 19 + 3 weeks of gestation. TTTS was resolved post-FLP and Doppler ultrasonography remained normal. The patient returned to her home country at 25 weeks of gestation and delivered eventually at 34 weeks of gestation via a Caesarean section for preeclampsia and worsening Doppler ultrasonography findings. Birth weights of the ex-donor and ex-recipient twins were 1,200 g and 2,000 g, respectively. As the patient had delivered at a foreign institution, there was no available data regarding neonatal outcomes, neonatal complications or total length of inpatient neonatal stay.

DISCUSSION
With the poor prognosis that accompanies TTTS in MC twin pregnancies, it is imperative for clinicians to achieve an early diagnosis so that early intervention may be possible and morbidity may be reduced. Over the last two decades, various interventions for the management of TTTS have been developed in a bid to improve the outcome of such pregnancies. These include serial amnioreduction, inter-twin septostomy, FLP and cord coagulation. All these treatment options have been found to result in improved survival rates compared to untreated TTTS. However, of the various treatment options, FLP (which was first described by De Lia et al117 in 1990) is thus far the only treatment aimed at treating the underlying pathophysiology. FLP has been shown to be a more effective first-line treatment than serial amnioreduction for severe TTTS diagnosed before 26 weeks of gestation, as demonstrated in the Eurofet study.15 Published series on FLP showed a 55%–73% survival rate, with a 4.2% neurological handicap rate.12-14 A Cochrane review in 2008 by Roberts et al15 also suggested that FLP of anastomotic vessels should be considered in the treatment of all stages of TTTS to improve perinatal and neonatal outcomes. However, the same review did not show that FLP reduces the risk of neurodevelopmental delay compared with other treatments.

Historically, serial amnioreduction has been one of the key therapies used in the management of TTTS. This intervention involves the repeated removal of large volumes of amniotic fluid, to improve outcome by reducing the risk of preterm labour and delivery secondary to polyhydramnios. It has also been postulated that serial amnioreduction improves fetal haemodynamics by decreasing pressure on the placental surface, thereby allowing increased blood flow to the uterus. Data from the International Amnioreduction Registry shows that at least one fetus survived to four weeks after birth in 71% of cases and both fetuses survived in 48% of cases.16 However, 25% of these survivors had abnormalities on neonatal cranial ultrasonography.16

Another therapeutic option for TTTS is cord coagulation using bipolar cautery or radiofrequency ablation, as a form of selective feticide. However, this therapy can only achieve a maximum survival rate of 50%, since the aim is to sacrifice one of the fetuses in a bid to achieve a better prognosis for the remaining twin. In one study of 14 patients, survival rates for the remaining twin after cord coagulation were 80%–90%, with no neurological abnormalities evident prior to discharge from hospital.17 However, this treatment may not be an acceptable option for parents of MC twins unless used as a last resort for pre-terminal TTTS, or if parents desire a better prognosis for one twin as opposed to a guarded prognosis for both.

One other intervention, inter-twin septostomy, which involves the intentional puncture of the inter-twin membrane, was first described by Saade et al to treat TTTS; the procedure has a survival rate of up to 83.3%.18 A randomised trial of septostomy versus amnioreduction has shown no difference in overall survival (70%); however, there was a decrease in the need for repeat procedures with septostomy (40% in the septostomy group vs. 70% in the amnioreduction group).19

Despite our study being a small case series owing to the small population base in Singapore, our results showed a 60% overall survival rate of pregnancies complicated by TTTS treated with FLP. Our results are comparable to those in other published reports.5,20,21 The severity of TTTS and gestation at delivery are the main factors in determining the perinatal outcome of such pregnancies.22 It is thus important to diagnose the condition as soon as possible in the second trimester and intervene when appropriate, in order to achieve the best possible outcomes for the MC twin pregnancy. In the present study, the median gestational age at diagnosis was relatively early at 19 weeks, with the median gestational age at intervention (i.e. FLP) at 20 weeks (i.e. a detection-to-intervention interval of only one week). The early diagnosis and intervention in our case series may have
contributed to the comparable overall survival rates of 60% found in our patients.

With regard to the technical aspects of FLP in our series, of the five patients, two had anterior placenta and the remaining three had posterior placenta. FLP procedures are known to be more technically challenging in the presence of an anterior placenta, and the approach of scope insertion may need to be altered accordingly to avoid injury and bleeding. For the two patients with anterior placenta, our approach included a detailed pre-fetoscopy ultrasonography mapping of the respective fetal positions, placental location and cord insertions, as well as a more lateral approach for the insertion of the fetoscope so as to avoid the anteriorly located placenta.

In the present study, there was one case of intrauterine death of the donor twin one day after FLP was performed, with the subsequent death of the ex-recipient twin 39 days post-FLP (Patient 2). One possible reason for the death of the donor twin is bleeding complication resulting from the procedure. Another possible reason for the death of the donor twin is unequal placenta sharing, which remained unchanged by the laser procedure. Therefore, after division of the communicating vessels, each fetus relies on the function of its own share of placental mass, which may be insufficient for some fetuses, as was possible in this case. On the other hand, the death of the ex-recipient twin was unlikely to be related to the FLP procedure, as the Doppler changes in the recipient twin had resolved post-FLP, and the ex-recipient twin died only later on Day 39 post-FLP for unidentified reasons.

In our series, single intrauterine death occurred in two of the five patients (Patients 1 and 4). However, there was no evidence of intrauterine demise or neurological handicap in the surviving twin after delivery in both patients. This is likely due to the success of the laser procedure in achieving division of all significant vascular communications between the twins, such that the death of one fetus no longer affected the haemodynamics of the remaining twin. This is one benefit of FLP compared to other available interventions, which do not focus on the separation of vascular communications. Other interventions, such as amnioreduction, do not target the underlying pathophysiology of TTTS and, therefore, the death of one twin would still subject the surviving twin to sudden changes in haemodynamics and an increased risk of intrauterine death or neurological handicap. A review of single intrauterine deaths in MC twin pregnancies showed only a 57% chance of having a healthy survivor.\(^{25}\)

The median gestational age at delivery in our series was 29 weeks. This is significantly earlier than most other reported large case series. For example, the median gestational age at delivery in the Eurofetus study was 33.3 weeks for FLP compared to 29 weeks for amniodrainage.\(^{20}\) The earlier median gestational age at delivery for the five patients in our centre may be attributable to a more cautious approach being taken in management in view of our limited experience with laser photocoagulation, as well as the commendable support extended by our hospital’s Neonatal Department, which has extensive experience in the management of severely preterm babies and an excellent track record of good neonatal outcomes.

Our study has several limitations. Firstly, the nature of the study, being a retrospective descriptive case series, did not allow us to compare the outcome of TTTS cases treated by FLP versus other methods such as amnioreduction or cord coagulation, which have similarly been used in our institution for the treatment of TTTS. Secondly, long-term neurological and developmental outcomes were not assessed in this case series. Another limitation was the small size of the study population, which is partly due to the small population of Singapore. Despite these limitations, the perinatal outcomes in our series appeared to be comparable to those reported internationally.\(^{24}\)

In conclusion, TTTS can be effectively treated by laser therapy with minimal maternal complications. However, management of this condition requires a specialised multidisciplinary team that is well-versed in the follow-up and detection of complications such as TTTS, and skilled in performing interventional therapies for the treatment of such complications. To achieve the best outcomes, such pregnancies should be managed by maternal fetal medicine specialists in specialised referral centres with tertiary neonatal facilities in order to handle the potential complications of prematurity, which are more likely to occur in these high-risk pregnancies.

REFERENCES


