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# Thrombosis after umbilical venous catheterisation: prospective study with serial ultrasound

Gerdina H Dubbink-Verheij <sup>1</sup>, Remco Visser,<sup>1</sup> Arno A Roest,<sup>2</sup> Cornelia Heleen van Ommen,<sup>3</sup> Arjan B te Pas,<sup>1</sup> Enrico Lopriore <sup>1</sup>

<sup>1</sup>Department of Neonatology, Leiden University Medical Center, Leiden, The Netherlands  
<sup>2</sup>Department of Pediatric Cardiology, Leiden University Medical Center, Leiden, The Netherlands

<sup>3</sup>Department of Pediatric Hematology, Erasmus MC Sophia, Rotterdam, The Netherlands

## Correspondence to

Mrs. Gerdina H Dubbink-Verheij, Neonatology, Leiden University Medical Center, Leiden 2333 ZA, The Netherlands; g.h.dubbink-verheij@lumc.nl

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## ABSTRACT

**Background** Umbilical venous catheters (UVCs) are associated with thrombus formation. Most studies on thrombosis in infants with UVCs focus on only one part of the route, and none assessed a control group of infants without UVCs.

**Objective** To determine the incidence and location of thrombi in infants after umbilical catheterisation and compare this with a control group of infants without umbilical catheters.

**Design** Prospective observational study with serial ultrasonography of the UVC route from the umbilico-portal confluence to the heart. Ultrasonography was performed until day 14 after catheterisation in cases and day 14 after birth in controls.

**Results** Thrombi in the UVC route were detected in 75% (30/40) of infants with UVCs in the study group, whereas no thrombi were detected in the control group of infants without UVCs (0/20) ( $p < 0.001$ ). Six thrombi (20%) were located in the right atrium. Most of these were also partly present in the ductus venosus. Six thrombi (20%) were located in the ductus venosus only, and in 12 infants (40%), the thrombus was at least partly located in the umbilico-portal confluence. Thrombi persisted after UVC removal in 25/30 cases. Two infants with thrombotic events were treated with low-molecular-weight heparin and resolution was found. In the other 23 infants managed expectantly, 2 died due to necrotising enterocolitis, 1 was lost to follow-up and in 20 spontaneous regression was seen.

**Conclusions** Thrombotic events occur frequently in infants after umbilical catheterisation. Most thrombi were asymptomatic and regressed spontaneously with expectant management. Routine screening for thrombi in UVCs is therefore not advised.

## INTRODUCTION

Umbilical catheters are frequently required for the management of critically ill infants. Umbilical venous catheters (UVCs) are used for intravenous administration of parenteral nutrition and medication. These catheters are relatively easy to insert and may be used for a longer period as compared with peripheral intravenous access. However, the advantages of UVCs should be balanced against the potential risks. Central venous catheterisation is reported to be the most common cause of neonatal thrombosis.<sup>1 2</sup> Catheter-associated thrombosis may be asymptomatic but may also result in thrombocytopenia, infection, pulmonary embolus, liver necrosis and, occasionally, even death.<sup>3–5</sup> Reported incidence of UVC-related thrombosis varies greatly from 2.2% to 43% due

## What is already known on this topic?

- ▶ Umbilical venous catheters are associated with thrombus formation.
- ▶ The incidence of thrombosis in infants without umbilical venous catheters is unclear.

## What this study adds?

- ▶ Most infants have thrombosis in the umbilical venous catheter route after umbilical venous catheterisation.
- ▶ Thrombosis in the umbilical venous catheter route in infants without umbilical venous catheters is not detected.
- ▶ Routine screening for thrombi in umbilical venous catheters does not seem necessary as most are asymptomatic and regress spontaneously.

to differences in study design and methodology.<sup>6–13</sup> UVCs are introduced in the umbilical vein to reach the umbilico-portal confluence and subsequently pass the ductus venosus to reach the junction between the inferior vena cava (IVC) and right atrium, which is the ideal location for the UVC tip.<sup>5 14</sup> The tip of UVCs is positioned too high when it is located in the right atrium or passing through the patent foramen ovale, within the left atrium or pulmonary vein.<sup>15</sup> Too low positioned UVCs have their tip in or below the umbilico-portal confluence or are malpositioned, for example, in the portal vein. Thrombus formation may occur in all parts of this UVC route. Most studies on thrombosis in infants with UVCs focus on only one part of the route, such as the liver or heart, not determining the incidence of thrombus formation in all locations together.<sup>6–11</sup> None of these studies investigated the occurrence of thrombosis in the UVC route in infants without UVCs.

The aim of this study was to determine the incidence and location of thrombi in infants after umbilical catheterisation with ultrasonography in the entire UVC route and to investigate if thrombosis in this route exists in infants without umbilical catheters.

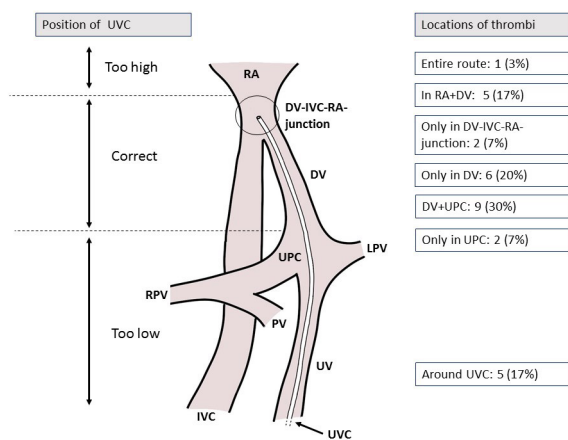
## METHODS

We conducted a prospective observational study at the neonatal intensive care unit of the Leiden University Medical Center, a tertiary care centre in the Netherlands. Written informed consent was obtained from the parents of participating infants.



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**Figure 1** Locations of the 30 detected thrombi, n (%). DV, ductus venosus; IVC, inferior vena cava; LPV, left portal vein; PV, portal vein; RA, right atrium; RPV, right portal vein; UPC, umbilico-portal confluence; UV, umbilical vein; UVC, umbilical venous catheter.

Infants of all gestational ages were considered for enrolment in the study group if a UVC was inserted. Infants were included from 1 October 2016 to 1 October 2017. We aimed for a convenience sample of 40 infants with and 20 without UVCs during this period. Umbilical catheterisation was performed according to local protocol. Estimated insertion length of the UVC was calculated based on the revised formula of Shukla.<sup>16</sup> An antero-posterior chest X-ray was routinely performed in all cases to determine the position of the catheter. Within an hour after catheterisation, the position of the catheter tip was determined by ultrasonography as well, and the presence and location of a thrombus was noted. Ultrasonography was performed by skilled neonatologists or paediatric cardiologists. Correct position of the UVC was defined as a UVC with the catheter tip in the ductus venosus or at the junction of the ductus venosus and IVC/right atrium by ultrasonography (see figure 1). Only infants with UVCs in the correct route with the catheter tip having passed the umbilico-portal confluence could be included in the study. Infants with UVCs positioned too low or with the catheter tip in blood vessels in the liver were excluded, because these UVCs were removed within 1 hour after insertion. These patients were also not eligible for the control group. UVCs that were positioned too high were pulled back to the correct position after ultrasonography. Ultrasound scans were repeated on day 3, 7 and 14 after catheterisation. Additional ultrasound scans were performed on the day of catheter removal. During the study period of 1 year, infants without an indication for a UVC were considered for enrolment in the control group. Matching cases and controls according to gestational age and birth weight was not possible as all infants below 28 weeks routinely receive umbilical catheters in our department. In the control group, ultrasonography was scheduled within 24 hours after birth, on day 7 and on day 14. In both groups in case of discharge of the patient before day 14, an ultrasound scan was performed within 24 hours preceding discharge. In case of non-availability of a skilled professional to perform ultrasonography, the examination was performed 1 or 2 days before or after the scheduled day.

Ultrasonography was performed with a Toshiba Aplio 400 or Aplio i700 machine (Toshiba Medical Systems Europe B.V., the Netherlands) with multifrequency transducers. Standard two-dimensional grey scale images were acquired from at least subcostal, abdominal and apical four-chamber views, sometimes

extended with parasternal short-axis views and were stored in digital format. The heart, IVC, ductus venosus and umbilico-portal confluence were visualised and screened for the presence of thrombi. A thrombus was defined as an echo dense structure within the heart or vessels or around the catheter observed in two dimensions. Except for thrombi in the ductus venosus, thrombi were classified as obstructive or non-obstructive based on the presence or absence of flow around the thrombus. Thrombi in the right atrium were classified as filling more or less than 50% of the atrium. Flow in the ductus venosus ceases within the first minutes to days independent of the presence or absence of a UVC.<sup>17</sup> Thus, flow was not used as a discriminating factor to assess thrombi in the ductus venosus.

Decisions to remove the UVC were made by the attending physician. Decisions to treat thrombi were based on the Dutch national guideline concerning catheter-related thrombosis in neonates.<sup>18</sup> This guideline advised a 'watch and wait' approach in most cases. If thrombi caused complete obstruction of blood flow or rapidly progressed without treatment, treatment with low-molecular-weight heparin (LMWH) was advised, as well as in thrombi filling >50% of the right atrium. Thrombolysis was only advised in cases with high risk of organ-threatening or life-threatening consequences.<sup>18</sup> Follow-up of thrombi after day 14 was performed if considered clinically indicated according to the attending physician.

All above data were recorded. Baseline characteristics of each infant, and clinical data, including gestational age, birth weight, duration of hospital admission, mortality, date and time of UVC placement and removal, timing of ultrasound scans, peripherally inserted central catheter insertion after UVC placement, position of catheter tip and presence and location of thrombosis, were extracted from the electronic patient dossier Metavision (iMD-soft, Leiden, The Netherlands). Presence and timing of necrotising enterocolitis stage 2a or higher,<sup>19</sup> culture-proven sepsis and thrombocytopenia (defined as a platelet count  $\leq 150 \times 10^9/L$ ) and treatment of thrombi were also recorded.

### Statistical analysis

Data are given as median with range unless otherwise mentioned. Incidences of thrombosis were compared with the  $\chi^2$  test. The Student's t-test for parametric and the Mann-Whitney U test for non-parametric data were used for comparisons of continuous variables. P values below 0.05 were considered significant. All statistical analyses were performed with IBM SPSS Statistics V.23.0.

## RESULTS

### Patient characteristics

We included 40 infants in the study group of infants with UVCs and 20 infants in the control group without UVCs. Patient characteristics in both groups are given in table 1.

Four patients in the study group died, due to necrotising enterocolitis (n=2) and pulmonary bleeding (n=2), and one patient died in the control-group due to necrotising enterocolitis (n=1).

### Incidence of thrombosis

Thrombi in the UVC route were detected in 75% (30/40) of infants in the study group, whereas no thrombi were detected in the control group (0/20) ( $p < 0.001$ ). Thrombi were first detected by ultrasonography at median day 6, in 14/30 cases (47%) at days 3–5, 12/30 (40%) at days 6–8 and 4/30 (13%) between day 9 and day 15. Thrombi were detected in 9/30 cases after

**Table 1** Patient characteristics of infants with UVC (study group) and infants without UVC (control group)

Variable	Study group (n=40)	Control group (n=20)	P value
Gestational age at birth (weeks)	27 (24–41)	29 (27–41)	0.06
Birth weight (grams)	1052 (600–3925)	1464 (1020–3300)	0.07
SGA, n (%)	6 (15)	0 (0)	0.07
Duration of hospital admission (days)	18 (3–102)	13 (3–44)	<b>0.04</b>
PICC received after UVC, n (%)	12 (30)	3 (15)	0.21
NEC, n (%)	5 (12.5)	1 (5)	0.36
Culture-proven sepsis, n (%)	11 (27.5)	2 (10)	0.12
Mortality, n (%)	4 (10)	1 (5)	0.51

Data are represented as median (range) unless otherwise specified. P values <0.05 were considered significant. Significant p values are depicted in bold. NEC, necrotising enterocolitis stage 2a or higher;<sup>19</sup> PICC, peripherally inserted central catheter; SGA, small-for-gestational-age, defined as a birth weight adjusted for gestational age below the 10<sup>th</sup> centile according to growth curves for Dutch neonates<sup>20</sup>; UVC, umbilical venous catheter.

UVC removal, in 4/30 cases at the day of UVC removal and in 17/30 cases before UVC removal. Thrombosis was not the reason for UVC removal in any case. Thrombi were detected in different parts of the UVC route (figure 1). Six thrombi (20%) were located in the right atrium. Five of these thrombi were also partly present in the ductus venosus, and one was located along the entire route including ductus venosus and umbilico-portal confluence as well. All right atrial thrombi filled less than half of the atrium and the two thrombi located at the junction of ductus venosus and IVC/right atrium were non-obstructive as well. Six thrombi (20%) were located in the ductus venosus only, and in 12 infants (40%) thrombi were at least partly located in the umbilico-portal confluence.

### Other aspects

Ultrasound data for study group and control group are given in table 2.

At initial ultrasound 5/40 UVCs (13%) were in correct position, 2/40 (5%) were placed with tips in the left atrium and 31/40 (78%) in the right atrium. In one UVC, the exact position of the tip could not be determined due to limited visualisation of the catheter by ultrasonography, and in one UVC, ultrasonography was not performed directly after catheterisation. Thrombus data of these cases are included in the analysis. Thrombocytopenia was seen in 6/40 infants with UVCs (15%) and in 2/20 controls (10%) (p=0.59) during admission. One infant with thrombocytopenia in the UVC group had no thrombus in the UVC route. In 3/5 infants with thrombocytopenia and a thrombus in the UVC route, the thrombocytopenia was observed before and in 2/5 after detection of the thrombus. In four cases, thrombocytopenia

**Table 2** Ultrasound data in study group and control group

	Study group (n=40)	Control group (n=20)
Follow-up duration after UVC insertion (days, median, range)	14 (2–16)	–
Follow-up duration after birth (days, median, range)	–	13 (1–17)
Position of UVC at initial ultrasound		
Correct	5 (13)	–
Too high	33 (83)	–
Not to determine	1 (3)	–

UVC, umbilical venous catheter.

was interpreted by the attending physician as a possible symptom of thrombosis. Two infants with thrombi had persistent catheter-related sepsis with *Staphylococcus aureus*. No other symptoms of thrombosis, such as loss of UVC patency, were noted.

### Follow-up of infants with catheter-related thrombi

Four of the 30 detected thrombi were visualised around the catheter and could not be visualised anymore after removal of the catheter. One infant with a thrombus around the UVC was lost to follow-up. The other 25 thrombi persisted after removal of the catheter. Two of 25 patients were treated with LMWH with resolution of the thrombus as a result. One, with thrombus formation in ductus venosus and umbilico-portal confluence, was treated because a thrombus in both venae iliacae communes developed after placement of another central venous catheter (CVC) and the other because progression of a thrombus in right atrium and ductus venosus during expectant management. All other patients were managed expectantly.

Two of the 25 patients with thrombi died during the neonatal period due to necrotising enterocolitis. In 12/25 patients, follow-up was considered not necessary, as the thrombi regressed spontaneously during the study period or were located in the ductus venosus (a non-functional structure after birth). The remaining 9/25 patients were discharged home without treatment with persisting thrombi in right atrium and/or umbilico-portal confluence. Thrombi in these nine cases were non-obstructive, filled less than 50% of the right atrium and were stable during hospital admission. One of these nine cases was lost to follow-up. At ultrasound follow-up in the outpatient department within 1 year after term age, the other eight thrombi all regressed and disappeared spontaneously.

### DISCUSSION

In this study, thrombosis in the UVC route was detected with screening by ultrasonography in 75% of infants after umbilical catheterisation, whereas no thrombi were demonstrated in the investigated location in infants without UVCs. This result emphasises that CVCs play an important role in the aetiology of neonatal thrombosis.<sup>20</sup> Studies described that thrombus formation in the UVC route could lead to thrombocytopenia, persistent sepsis, liver damage, portal hypertension, symptoms of right heart failure and pulmonary embolism.<sup>21–23</sup>

We prospectively screened the complete route of the UVC from the umbilico-portal confluence up to and including the heart, whereas previous studies mainly focused on one or two parts of this route. This might explain the higher incidence of UVC-related thrombosis (75%) when compared with previous reports (2.2%–43%).<sup>6–13</sup> The incidence of thrombi located only in the umbilico-portal confluence in this study was 30% (12/40), which is comparable with the incidence reported in previous studies. However, the incidence of thrombi located in the right atrium, IVC and ductus venosus was much higher, 58% (23/40) when compared with previous reports (10%–30%).<sup>6–8</sup> The use of novel ultrasound machines with higher sensitivity to detect thrombi compared with the equipment used in the past may possibly explain the higher incidence of thrombi compared with older studies. Methodological differences related to the use of different ultrasound screening programmes or differences in the screened population may be an explanation as well. Suggested risk factors for catheter-related thrombosis, such as being small-for-gestational age, dwelling time of the UVC and the use of parenteral nutrition could differ in our population compared with others and be of influence on the incidence of thrombosis.<sup>22</sup>

However, the sample size of our group is too small to confirm the influence of risk factors.

Neonatal thrombosis is in 89%–94% of cases catheter related, but reliable data on thrombosis in infants without catheters, especially in the UVC route, are not available.<sup>2 20</sup> The ductus venosus is a fetal structure that after birth closes permanent with fibrotic transformation into the ligamentum venosum. Some authors suggest thrombus formation contributes to this process as well.<sup>17 24</sup> However, we did not observe this in our study. Hence, although our control group was just a small group of infants, the thrombotic incidence of 0% in this group is an important finding.

Neonates are at increased risk of thrombosis due to small vessel diameters and disruption of haemostatic balance by numerous acquired and prothrombotic disorders.<sup>5 21 22 25</sup> Intravascular catheters can cause thrombosis by damaging the epithelium and introducing a foreign thrombogenic surface.<sup>5</sup> In clinically unstable infants at increased risk of thrombosis, CVCs are needed to provide optimal neonatal care. The umbilical vein was for decades an obvious possibility for catheterisation and, in many cases, the first choice to get intravenous access in infants. The high incidence of thrombosis in infants with UVCs demonstrated in this study suggest that our policy on use of UVCs needs to be reconsidered. However, CVCs are frequently needed in sick infants and other types of central catheters than UVCs are associated with thrombosis as well. Replacement of the UVC by another type of catheter may not necessarily lead to less complications.<sup>6 13 22 26 27</sup>

Management of UVC-related thrombi is based on expert opinion guidelines and the impact of these thrombi on outcome is not clear.<sup>20–22 26 28</sup> As shown in our study, spontaneous resolution of UVC-related thrombi detected by screening is likely to occur. Therefore, a ‘watch and wait’ approach in asymptomatic thrombi detected by screening with ultrasonography may be justified.<sup>26</sup> However, it is unclear how many infants with catheter-related thrombi will develop long-term consequences later in childhood. To determine the natural history, and possible consequences of catheter-related thrombosis, a larger prospective study is needed. Currently, a prospective study in neonates with thrombi is being performed in the Netherlands (the Neoclot study).<sup>18</sup> Because clinical implications of thrombi are not clear yet, and most thrombi detected by screening disappear spontaneously, routine screening apart from research settings cannot be advised. This is also confirmed by Haddad *et al*, who report that the majority of catheter-related thrombi they detected was stable and detection led infrequently to changes in patient management.<sup>28</sup>

Care should be taken when interpreting the results of our study due to several limitations, including the relatively small number of patients. In addition, we excluded infants with UVCs positioned too low or malpositioned with catheter tips in veins in the liver, because these catheters were removed shortly after catheterisation and not used. It would have been interesting to know the incidence of thrombi in this separate group, with catheters being in situ for a very short period.

In conclusion, our study demonstrates that thrombi are frequently detected after umbilical catheterisation by ultrasonography. Most thrombi are asymptomatic and disappear without treatment. Thus, routine screening for thrombi in UVCs is not advised. However, given the high risk of thrombus formation in UVCs, these catheters should be used with caution and removed promptly if the clinical indication is no longer present.

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**Contributors** GHD-V was the executive researcher of this study. She performed literature search, data collection, data analysis, data interpretation, writing and

submitting of the report. RV was involved in study design, data collection and writing of the report. AAR was involved in study design, data collection and critical revision of the content of the report. ChVO was involved in writing and critical revision of the content of the report. ABtP was involved in study design, data collection and interpretation and critical revision of the content of the report. EL was the project leader and performed literature search, coordinated data analysis, data interpretation and writing and editing of the report. All authors gave approval for the final version to be published and agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of a part of the work are appropriately investigated and resolved. No honorarium, grant or other form of payment was given to anyone to produce the manuscript.

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#### ORCID iDs

Gerdina H Dubbink-Verheij <http://orcid.org/0000-0003-0734-2233>  
Enrico Lopriore <http://orcid.org/0000-0002-3513-5066>

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