Cerebral Venous Thrombosis: What’s New?

Diana Aguiar de Sousa¹,²,³

¹ Department of Neurosciences and Mental Health (Neurology), Hospital Santa Maria - CHULN, University of Lisbon, Lisbon, Portugal
² Faculdade de Medicina, Universidade de, Lisboa, Portugal
³ Instituto de Medicina Molecular João Lobo Antunes, Lisbon, Portugal

Address for correspondence: Diana Aguiar de Sousa, Department of Neurosciences and Mental Health (Neurology), Hospital Santa Maria, University of Lisbon, Av. Prof. Egas Moniz, 1649-028-Lisbon, Portugal (e-mail: dianasousa@campus.ul.pt).

Abstract

Keywords
► cerebral venous thrombosis
► brain lesion
► recanalization
► risk factor
► anticoagulation

Thrombosis of the cerebral veins and sinuses (CVT) is a distinct cerebrovascular disorder that, unlike arterial stroke, most often affects children and young adults, especially women. In this review, we will summarize recent advances on the knowledge of patients with CVT.

Epidemiology

Cerebral venous thrombosis (CVT) is a less common form of stroke. In the largest published prospective multinational cohort of patients with CVT—the International Study on Cerebral Vein and Dural Sinus Thrombosis (ISCVT),¹ the proportion of females was 75% and the median age at the thrombotic event was in the fourth decade of life. Other cohorts, from all over the world, found similar trends,² with most adults with CVT being aged 20 to 50 years and less than 10% older than 65 years.³ Importantly, the female predominance is restricted to the fertile age, as it is not seen in pediatric cohorts or in elderly patients.⁴ The most accurate estimates concerning incidence of CVT come from high-income countries, where two studies conducted with this aim indicated an overall prevalence of 13 to 16 cases per million per year.⁵ Also, a recent retrospective cohort study from the United States reported an incidence between 14 and 20 cases per million population (20–27 cases per million in females).⁶ Of note, this study also described an increase in incidence over time in older females and males, and a higher incidence rate in black Americans.

Clinical and Imaging Manifestations

Knowledge on the presentation and prognosis of CVT was very much improved in the last two decades due to the more widespread availability of noninvasive diagnostic techniques and the publication of the first large cohort studies. CVT has a more varied clinical presentation than other stroke types and may be difficult to recognize, which may contribute to underdiagnosis.⁷ The clinical onset is more often subacute, but an acute presentation (<48 hours) is also observed in about one-third of cases and a more chronic course is also possible, although it is less frequent.³ Symptoms and signs of CVT can be grouped in presenting syndromes, of which the most common are isolated intracranial hypertension syndrome, focal syndrome, and encephalopathy.⁷

The spectrum of parenchymal damage due to intracranial venous thrombosis includes a wide range of manifestations, from mild vasogenic edema to extensive parenchymal hemorrhage. Parenchymal brain lesions in CVT are often classified as nonhemorrhagic and hemorrhagic.⁸ These changes are better depicted in magnetic resonance imaging (MRI), which has increased sensitivity for the identification of parenchymal changes, especially the nonhemorrhagic, and also allows distinction between vasogenic and cytotoxic edema. In the ISCVT cohort, 64% of patients had parenchymal changes on computed tomography or MR.³ About half of the cases have a hemorrhagic component that often expands to the cortical surface, which may help in differentiating from the typical localization of parenchymal hypertensive bleeds. Juxtacortical flame-shaped lobar hemorrhages in the parasagittal frontal and parietal lobes, between the superficial
and deep venous drainage systems, were described as a very specific finding in patients with CVT, particularly those with superior sagittal sinus thrombosis.\textsuperscript{9} The Brush sign, an MRI finding previously described in patients with ischemic stroke, can also be found in patients with CVT, in whom it is associated with ipsilateral parenchymal brain lesion, larger extent of thrombosis, and more severe clinical presentation.\textsuperscript{10} It corresponds to an abnormally accentuated hypointensity of the subependymal and medullary veins that can be seen in paramagnetic sensitive sequences (T2* or susceptibility-weighted imaging) and is a likely marker of hypoperfusion and increased collateral circulation.

**Pathophysiology of Brain Damage**

Despite the progress made in the last years, significant evidence gaps persist in the understanding of the pathophysiology of brain damage and the mechanisms underlying the benefit of the available treatment strategies. Although there are several anecdotal reports of tissue recovery after recanalization, particularly following endovascular treatment, the association between early venous recanalization and tissue fate was not established until recently. In a prospective cohort study of patients treated with anticoagulation, we found that patients with persistent venous occlusion by day 8 after starting of anticoagulation treatment were more likely to have early worsening of nonhemorrhagic brain lesions.\textsuperscript{13} Importantly, patients achieving venous recanalization within the first week of therapeutic anticoagulation had a higher chance to show early regression of the nonhemorrhagic parenchymal lesions, comparing with patients with persistent occlusion. Besides suggesting that vessel recanalization is relevant for the evolution of brain damage, this finding raises the hypothesis that anticoagulation-induced early partial recanalization contributes to the clinical improvement often seen in these patients just a few days after treatment start. Contrary to what is described in ischemic stroke, lesions showing diffusion restriction were fully reversible in most cases, particularly in patients achieving early venous recanalization.\textsuperscript{11} This suggests that, in CVT, what was thought to represent cytotoxic edema can still represent viable brain tissue, and may imply that a longer time window for treatment options exists in CVT, even in patients showing the so-called venous infarction.

**Risk Factors**

The temporal profile of risk factors is a crucial feature because persistent predisposing conditions usually have specific implications on the long-term risk of recurrent venous thrombotic events. Therefore, risk factors are usually classified in permanent/nonmodifiable, such as genetic thrombophilia, and precipitant/transient, such as pregnancy or infection.\textsuperscript{12}

CVT mostly affects young women, as the most common risk factors are female-specific, particularly pregnancy and hormonal contraception. Currently, oral contraceptives are the most frequent risk factor for CVT in women in Western countries.\textsuperscript{1} The overall risk of CVT associated with use of oral contraceptives was estimated to be seven times that of nonusers.\textsuperscript{13} However, the risk will be particularly high among those who carry other risk factors, such as obesity or hereditary thrombophilias.\textsuperscript{14,15} In fact, multiple risk factors are found in about half of patients.\textsuperscript{1} With regard to pregnancy and puerperium, the risk of any venous thrombosis was estimated to be increased by 4- to 5-fold during pregnancy and by 22-fold during the first 6 weeks after delivery, compared with non-pregnant women.\textsuperscript{16,17} Like for other venous thrombotic events, the risk of pregnancy-related CVT is higher in third trimester and puerperium, especially in the first 6 weeks postpartum, as confirmed by a recent case-control study conducted in high-income countries.\textsuperscript{18} Anemia and obesity have been also added to the list of risk factors for CVT over the past few years.\textsuperscript{15,19} Recently, there has been increased attention on the possible increased risk of CVT in patients with SARS-CoV-2 infection, as described for other thrombotic complications.\textsuperscript{20} Although the number of cases described so far is small\textsuperscript{21} underdiagnosis is likely, especially in patients with severe COVID-19. Further studies are warranted to confirm whether SARS-CoV-2 infection is a new risk factor for CVT.

With regard to the so-called permanent risk factors, the most common predisposing conditions are genetic prothrombotic disorders.\textsuperscript{1} Several publications from different regions of the world demonstrated that inherited thrombophilia increases the risk of CVT, especially in children.\textsuperscript{22} Acquired prothrombotic disorders are also found in a significant proportion of CVT cases, such as antiphospholipid syndrome (6–17% of cases).\textsuperscript{23} Of note, there are several reports of CVT as a presenting manifestation of this syndrome.\textsuperscript{22} Cancer has been confirmed to be a powerful risk factor for the development of CVT, as the risk was estimated to be increased fivefold in patients with history of cancer.\textsuperscript{24} However, in patients with hematological neoplasms, the risk of CVT can be particularly increased, especially in the first year after diagnosis, with an estimated odds ratio for CVT of 86, according to a recent case-control study.\textsuperscript{25} Some specific forms of hematological cancer are more often associated with CVT, namely acute lymphoblastic leukemia\textsuperscript{26} and polycythemia vera.\textsuperscript{20} Specific hematopoietic agents, such as L-asparaginase, and other iatrogenic factors, such as lumbar punctures for intrathecal therapy, can also precipitate CVT. Several inflammatory conditions are also associated with increased risk of CVT. Behçet disease, is often complicated by CVT, with an estimated incidence of 3 cases per 1,000 person-years,\textsuperscript{27} corresponding to approximately 200 times the risk in the general population. In about 30% of these cases, CVT occurs in patients without a previous diagnosis of Behçet disease.\textsuperscript{27}

Although the most common risk factors for CVT correspond to systemic disturbances, local conditions have also an established association with CVT, including head trauma,\textsuperscript{28} neurosurgical procedures,\textsuperscript{29} and infections, especially of the head and neck.\textsuperscript{30} Currently, these risk factors contribute to a smaller proportion of cases, despite considerable geographical variations in prevalence. Infectious causes, in particular, have substantially declined and septic CVT is uncommon nowadays, as a result of early treatment with antibiotics. Still, in
contemporary multinational cohorts, infection was associated with 8 to 11% of cases of CVT. No underlying etiology or risk factor for CVT is found in approximately 15% of adult patients.

**Venous Recanalization**

Considering data from previously published cohorts that included more than 800 patients treated with anticoagulation, a pooled proportion of 85% of cases showed recanalization of at least one of the thrombosed vessels during follow-up (95% confidence interval [CI], 80–89; \( P = 58\% \)). However, the methods and time points for evaluation varied across studies and there was significant heterogeneity in the results. To assess whether venous recanalization could occur earlier, even in patients treated only with anticoagulation, at what could be a more critical time window for the association with imaging and clinical outcomes, we have recently conducted a prospective study in which we were able to show that recanalization starts early in patients receiving therapeutic anticoagulation, as three quarters of patients had no persistent venous occlusion at day 8. However, venous recanalization progresses with time, and complete recanalization was only reached in about half of the patients at 90 days. Younger age was a predictor of early recanalization and there was a trend to an increased rate of early recanalization in patients showing the susceptibility vessel sign at admission. Although an association between persistent venous occlusion and worse functional outcome was found in the meta-analysis of previous cohort studies, that was not confirmed in this prospective cohort study with serial imaging at specific time points. Even so, the most severe presentations tended to be underrepresented and the sample size was likely to be underpowered to detect an association with long-term functional outcome, in a disease that most often has a favorable prognosis, as measured by the Rankin scale. Lack of complete recanalization was not a predictor of persistent headache.

**Treatment**

The European guidelines on the management of CVT have been updated in 2017. However, the quality of evidence was classified as low or very low for most of the recommendations, as they are mostly based on data from observational studies. Therapeutic anticoagulation is the standard treatment in CVT. The recommended antithrombotic treatment in the acute phase is heparin, either intravenous nonfractionated heparin or subcutaneous low-molecular-weight heparin (LMWH). Two trials and one observational study showed a trend in favor of LMWH in mortality and functional outcomes. However, these studies had several methodological limitations, reducing the confidence on the conclusions.

Based on the hypothesis that rapid venous recanalization can prevent further deterioration and improve prognosis in severe CVT, use of endovascular treatment has been reported since the 1980s in hundreds of case reports and case series. The main endovascular approaches for the treatment of CVT are an intravascular chemical thrombolytic, mechanical thrombectomy, or the combined approach, using both techniques. A 2017 systematic review of case series with at least three cases treated with endovascular treatment included 235 patients. In this analysis, the authors found a mortality of 14% among patients receiving endovascular treatment, worsening or new intracranial hemorrhage in 9%, and complete recovery in 35%. Patients selected to the intervention were mostly severe cases, with 40% having encephalopathy or coma. The rate of complete recanalization was 69%. A more recent systematic review that included 393 patients found that the combination of mechanical thrombectomy with local thrombolysis, when consistently performed across the cohort, tended to be associated with better outcomes. However, without a control group, no firm conclusions on the efficacy and safety of thrombectomy can be inferred from these analyses. Still, an evaluation of the United States Nationwide Inpatient Sample between 2004 and 2014 estimated that approximately 3% of all CVT patients received endovascular treatment (1,248 patients). There was high heterogeneity in the used techniques and no information was available on the recanalization rate. Due to the very low quality of available evidence, the European guidelines do not make any recommendation, and suggest not using endovascular treatment in acute CVT patients with a pretreatment low risk of poor outcome. Recently, the neutral results of the first randomized comparison of adjunctive EVT versus standard treatment with anticoagulation have further increased the uncertainty on the effectiveness of this intervention in patients with severe CVT. The TO-ACT trial included patients having at least a predictor of poor outcome and decisions regarding the protocol for the endovascular intervention were left to the treating interventionalist. The trial was terminated prematurely for futility, after inclusion of 67 patients, and no difference in clinical outcome was detected between those allocated to anticoagulation only or adjunctive endovascular treatment (modified Rankin’s scale of 0–1 in 67% vs. 68% patients, in the intervention and standard treatment arm, respectively). Perforation of sinus or cortical vein was reported in 3 three cases (9%), although this was not associated with symptomatic bleeding. Further research on the pathophysiology of brain lesions, prognostic markers, and intervention techniques directed to the venous system should contribute to improve patient selection, as well as the safety and efficacy of these neurovascular interventions in CVT. Herniation due to unilateral mass effect is the major cause of death in CVT. Although the evidence for the use of decompressive surgery in CVT comes from uncontrolled studies, it consistently suggests patients with impending brain herniation are best treated with decompressive surgery. Particularly, several single-center series and one multicenter registry and systematic review of individual patients showed that decompressive surgery is lifesaving and often results in good functional outcome. A prospective register of decompressive surgery in CVT (DECOMPRESS-2) that included more than 100 patients has been recently finished and results are expected soon.

Treatment of the associated conditions/risk factors and management of symptoms and complications is also an
important part of the management of patients with CVT. Acute symptomatic seizures are a common manifestation, occurring in about one-third of patients.45 Late seizures occur in 11% of patients, especially in patients with intracranial bleeding, acute symptomatic seizures, or who underwent decompressive hemicraniectomy.46

Long-Term Management

After the initial period of parenteral anticoagulation, oral anticoagulants should be started, unless there is a specific contraindication. Recently, an exploratory randomized controlled trial (RE-SPECT CVT) comparing the efficacy and safety of dabigatran 150 mg, bid, versus dose-adjusted warfarin (INR 2–3) during 6 months, was completed, after allocating 60 CVT patients to each treatment arm.47 Oral anticoagulation was started 5 to 15 days after the beginning of parenteral anticoagulation, in clinically stable patients. The risk of recurrent venous thrombotic events was low in both arms, with similar rates of bleeding, suggesting that both dabigatran and warfarin may be safe and effective for preventing recurrent thrombotic events in patients with CVT. There is an ongoing open-label phase II trial assessing the safety of rivaroxaban compared with warfarin in patients with CVT (SECRET trial NCT03178864). Contrary to the previous trial, oral anticoagulation can be started in the acute phase, instead of parenteral anticoagulation. The EINSTEIN Junior study is also evaluating the efficacy and safety of rivaroxaban in children with venous thrombosis, including CVT (NCT02234843).

Based on the largest cohorts studies with long-term follow-up, the risk of recurrent events in patients with previous CVT is estimated to be around 2 events per 100 patient-years for recurrent CVT and approximately 4 events per 100 patient-years for noncerebral venous thrombosis,2,48 with most events occurring after anticoagulant therapy withdrawal. Importantly, the rate of recurrence seems to persist with time, as demonstrated in a long-term French cohort study including 187 patients, in which 18% of patients had a recurrent thrombotic event in the 10 years that followed CVT.49 In adults, risk factors for recurrence of CVT are male gender, myeloproliferative diseases (polycythemia vera or essential thrombocytopenia), prior venous thrombotic events, and severe thrombophilia.50–52

Oral anticoagulation after the acute phase of CVT contributes to the prevention of further venous thrombotic events, including recurrence of CVT. However, there have been no randomized controlled trials or prospective controlled studies assessing optimal duration of oral anticoagulation in this setting.32 The first trial that addresses this question (EXTending Oral Anticoagulant treatment after acute Cerebral Vein Thrombosis, EXCOA-CVT) is currently ongoing.53 Meanwhile, oral anticoagulation is usually maintained for 3 to 12 months after acute CVT.32 In patients with genetic or acquired prothrombotic conditions (e.g., antithrombin deficiency, antiphospholipid syndrome, and active cancer) at high risk of recurrent venous thrombosis, or with history of prior venous thrombotic events, lifelong anticoagulation should be considered.30

Future health conditions associated with increased risk of recurrent thrombotic events should also be a concern. As CVT is more common in women at fertile age and thromboembolism is a leading cause of maternal mortality, safety of future pregnancies is a common question in clinical practice. Specifically, besides counseling, a decision has to be made on whether to offer antithrombotic prophylaxis or not during pregnancy and/or puerperium. The currently available evidence, which was substantially increased in the last years by new follow-up studies54 and systematic reviews,55,56 supports medical counseling in favor of not contraindicating future pregnancy in women with previous CVT, although they should be informed on the increased risks.32 Also, in women with a history of CVT, heparin prophylaxis was associated with lower rates of pregnancy-related recurrent thrombotic events and a significantly lower risk of miscarriage.54–56

Prognosis

Overall, death or dependence occurs in approximately 15% of patients, despite medical treatment.1 However, mortality rates above 30% have been reported in case series of patients with severe CVT admitted to an intensive care unit.57,58 mostly due to large herniating brain lesions.

The clinical course of CVT is unpredictable in the first days after diagnosis, and about one quarter of patients deteriorate in the acute phase.1 Among the surviving patients, approximately 6 to 10% have severe and permanent disability.1 Predictors of poor outcome include older age, male sex, coma, mental status disorder, intracerebral hemorrhage, thrombosis of the deep venous system, infection of the central nervous system, cancer, and hyperglycemia at admission.1,59

In recent years, there was increasing interest in the subtle consequences of CVT in patients with apparently full recovery. Even though approximately 80% of patients recover from CVT without physical disability, many of these patients experience residual chronic symptoms. About half of patients report headache during follow-up, and severe headaches that require bed rest or hospital admission persist in 14% of patients.1 More than half of survivors of CVT report subtle neuropsychological difficulties or depression. These complaints are often associated with a negative effect on employment status, and 20 to 40% of patients are unable to return to their prior working life.60

Conflict of Interest

Dr. Aguiar de Sousa reports nonfinancial support from Boehringer Ingelheim, outside the submitted work.

References


Hämostaseologie  Vol. 41  No. 1/2021  © 2021. Thieme. All rights reserved.


Ferro JM, Crassard I, Coutinho JM, et al;Second International Study on Cerebral Vein and Dural Sinus Thrombosis (ISCVT 2)