Evidence-based consensus on the insertion of central venous access devices: definition of minimal requirements for training


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Editor’s key points

• This review presents consensus on standard minimal requirements for training on central venous access devices.
• An international task force generated an evidence-based consensus.
• The task force proposed 16 recommendations.
• Standardized education, simulation practice, and supervised insertions are the key to ensuring safe and competent practice.

Summary. There is a lack of standard minimal requirements for the training of insertion techniques and maintenance of central venous access devices (CVADs). An international evidence-based consensus task force was established through the World Congress of Vascular Access (WoCoVA) to provide definitions and recommendations for training and insertion of CVADs. Medical literature published from February 1971 to April 2012 regarding ‘central vascular access’, ‘training’, ‘competency’, ‘simulation’, and ‘ultrasound’ was reviewed on Pubmed, BioMed Central, ScienceDirect, and Scopus databases. The GRADE and the GRADE-RAND methods were utilized to develop recommendations. Out of 156 papers initially identified, 83 papers described training for central vascular access placement. Sixteen recommendations are proposed by this task force, each with an evidence level, degree of consensus, and recommendation grade. These recommendations suggest central venous access education include didactic or web-based teaching with insertion procedure, infection prevention, complications, care, and maintenance of devices, along with laboratory models and tools for simulation practice incorporating ultrasound. Clinical competence should be determined by observation during clinical practice using a global rating scale rather than by the number of procedures performed. Ensuring safe insertion and management of central venous devices requires standardized education, simulation practice, and supervised insertions.

Keywords: catheter-related infections, prevention and control; catheterization; central venous access; central venous, standards; clinical competence; competency; complications; computer-assisted instruction; consensus; evidence-based medicine; education; GRADE; guideline; humans; infection; internship and residency; programme development; programme evaluation; RAND; supervision; simulation; subclavian vein; training; ultrasound guidance; ultrasonography; vascular access; vascular surgical procedures

Education surrounding the insertion of central venous access devices (CVADs) remains undefined. Training is defined as the acquisition of knowledge, skills, and competence related to a specific activity or procedure. Understanding and establishing the level of education required for safe insertion procedures and management of CVADs is the focus of this publication. There is variability of knowledge and competency among inserters which is represented quantitatively by the number of complications that occur from patient to patient.1 It has been demonstrated that a systematic training process, including ultrasound instruction before patient insertions, reduces mechanical and infectious complications.2–6

Current CVAD literature related to training, supervision, and competence acquisition does not define a fully standardized programme for trainees; nor does it establish guidelines for supervisors. No standard didactic or simulation training is currently required before the insertion of CVADs by clinicians in training other than supervision of an unspecified number of insertions. The supervision requirements do not specify the role, experience, or competence of the supervisor.
Healthcare workers involved in the placement of CVADs using ultrasound guidance need appropriate education and training to ensure patient safety and avoid major complications with the insertion of CVADs. Basic knowledge of anatomy, ultrasound physics and imaging, and infection prevention strategies have been proposed for the standard didactic education. These recommended topics are necessary for adequate understanding and safety of the insertion procedure.

There are two areas of focus to be addressed in any CVAD educational course; insertion and management. The insertion method and site selected affect the amount of risk involved related to trauma, colonization, and the ability to complete therapy successfully. Even the decision to choose a particular type or size of device contributes to the risk for infection and the development of thrombosis. Furthermore, there is a synergistic effect in which risk factors for one event may impact the incidence of other complications. For example, there is a direct association between catheter-related thrombosis and infection; the incidence of thrombosis increases with multiple insertion attempts which then increases the risk of infection.

A growing body of knowledge points to simulation training as a key to safe patient insertions by advocating competency-based education and multidisciplinary practice models. Application of ultrasound guidance with CVAD insertions reduces insertion-related complications, increases success, and establishes a process for vascular access based on safety and vein preservation. The safety afforded with ultrasound-guided insertions dictates that this technique be included in the educational process of any central venous device placement. Educational processes and supervised insertions are needed for healthcare providers to establish credentialling for CVAD procedures in any healthcare facility.

In keeping with recommendations and guidelines, standard education on principles of insertion and infection prevention practices should be provided to all CVAD inserters initially and at least annually.

To address the issue of standardization of CVAD training, a task force was formed by the World Congress of Vascular Access (WoCoVA) with a goal to create evidence-based recommendations for minimal education and training for central venous device insertion and management.

Methods

Eight worldwide educational course experts on vascular access device placement, not supported by industry, were identified by WoCoVa in 2010 to create an evidence-based consensus on minimal requirements in training in central venous device placement. These experts qualified based on a minimum authorship of two peer-reviewed articles published in the past 10 yr related to this topic, and additional activities including teaching and speaking on vascular access. Seven panellists accepted who then created a roadmap to achieve a final document with evidence-based recommendations on CVAD education. A search of medical literature was performed using two methods to avoid selection bias: the first method entailed a systematic search by all panel experts. Medical subject headings including ‘central vascular access’, ‘training’, ‘competency’, ‘simulation’, ‘infection’, ‘complications’, and ‘ultrasound’ were searched on Pubmed, BioMed Central, ScienceDirect, and Scopus databases including articles dated from February 1971 to April 2012. A professional medical librarian from the National Neurological Institute Besta in Milan supplemented this first search with a hand search based on selected articles from the expert panel. The second method entailed a systematic search of English language articles from the same period by an epidemiologist (M.E.) assisted by a professional librarian. The two bibliographies were then compared for thoroughness and consistency. Out of 156 papers initially identified, 83 papers were linked with training in vascular access. The GRADE and the GRADE-RAND methods were utilized to develop the 16 recommendations. The GRADE method utilized two phases in the development of these evidence-based recommendations. This methodology has been previously detailed in the published literature. There are 15 factors that are typically considered in the GRADE process. The level of evidence quality was scored according to nine factors. The final classification of evidence quality was divided into three levels (A, high; B, moderate; C, low). The transformation of evidence into a recommendation was a function of the panel evaluation of five factors. The GRADE system has not standardized this decision-making process of the expert panel. In an effort to standardize this evidence processing, the methodology committee of this working group selected the Rand Appropriateness Method (RAM). The panellists held conference calls in which they discussed the topics of the Consensus and voted separately on all recommendations using a web-based voting system. The voting process required expert decisions utilizing GRADE factors such as outcome importance and evidence-to-recommendation transformers. This process provided a structured and validated method for expert panel activities. In addition, it standardized statistical methodology for determining the degree of agreement to serve as a foundation for deciding about the recommendation grade (weak vs strong).

Results

Eighty-three articles were analysed and voted upon according to the GRADE factors. Sixteen recommendations were proposed, each with an evidence level, degree of consensus, and recommendation grade (Table 1).

Adult learning methods

These teaching methods follow a consistent scientific approach or educational style to engage the student’s mind. The approach to teaching and learning with regard to CVAD insertion should be underpinned by a constructivism or adult learning philosophical framework such as experiential learning.
<table>
<thead>
<tr>
<th>Section code</th>
<th>Recommendation</th>
<th>Level of evidence</th>
<th>Degree of consensus, strength of recommendation</th>
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<tbody>
<tr>
<td>W_R1</td>
<td>Didactic Educational Content Education on anatomy and physiology is relevant to ultrasound guidance, vein assessment, and selection, and complications that may occur at any possible CVAD insertion site. Specific knowledge of vessel characteristics, skin, peripheral nervous, respiratory, and cardiac systems is included. Insertion sites for central venous access include: upper arm, chest, neck, and inguinal. Anatomy and physiology include both typical and variant presentations.</td>
<td>C</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R2</td>
<td>Didactic Education on Ultrasound-guided Technique Education for using ultrasound for vein access and assessment is a component of any CVAD insertion course. Education includes: physics of ultrasound, image optimization, image analysis, anatomical assessment of both normal and variant anatomy, and simulation skills training. Landmark techniques are also components of CVAD education programmes in case circumstances arise that do not permit the use of ultrasound. Educational programmes emphasize the indisputable benefit of ultrasound for CVAD insertion.</td>
<td>B</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R3</td>
<td>Education on Tip Placement Education on CVAD tip placement is based on three essential features that reduce risk: high blood flow, parallel positioning to the vessel, and pulsatility of blood flow. To achieve these three ideal factors of tip location, the tip of the CVAD is at or near the cavo-atrial junction.</td>
<td>C</td>
<td>Very good, strong</td>
</tr>
<tr>
<td>W_R4</td>
<td>Infection Education to prevent catheter-related primary bloodstream infections focus on pre, intra, and post-procedure insertion, care, and maintenance. Three main sources of infection contamination: the catheter–skin junction (extraluminal), the needleless connector and hub (intraluminal), and iatrogenic contamination. Topics to include: site selection, device selection, indications for use, maximal barrier precautions, hand washing, skin antisepsis, and the use of central line checklists are included in CVAD insertion course. Observers, empowered to stop a procedure if contaminated, are part of the implementation of the educational programme.</td>
<td>A</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R5</td>
<td>Education on Catheters Topics in educational programmes include catheter: material composition, size, features, and number of lumen. Focus education on various risks vs benefits of catheter features for both short- and long-term use.</td>
<td>C</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R6</td>
<td>Education on Procedure Any central venous access educational programme includes insertion procedure for respective device placement, with both typical and variant scenarios.</td>
<td>B</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R7</td>
<td>Education on Care and Maintenance CVAD care and maintenance is addressed in CVAD education programme. Complications from CVAD use are related to the insertion procedure or from continued care and maintenance of the device.</td>
<td>B</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R8</td>
<td>Instructors Only providers with demonstrated qualification and competency can provide education on CVAD insertion. The basic tenants of qualification and competency are addressed within the content of a CVAD education programme.</td>
<td>C</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R9</td>
<td>Ultrasound Simulation A simulation practice on ultrasound anatomy is included in the curriculum of CVAD placement training.</td>
<td>B</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R10</td>
<td>Anatomical Models Inanimate models/simulators replace the human anatomy as clinicians follow minimal requirements to allow for the detection of main vessels and surrounding structures</td>
<td>A</td>
<td>Perfect consensus, strong</td>
</tr>
<tr>
<td>W_R11</td>
<td>Objective Grading of Proficiency Levels of performance of theoretical and practical skills are objective and the trainee completes every step of the final assessment in order to obtain the certification of proficiency in ultrasound CVAD positioning.</td>
<td>C</td>
<td>Perfect consensus, strong</td>
</tr>
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Continued
Experiential learning is a model of learning, in which learning/meaning is derived from direct experience. According to Bruner, professional judgement and higher level skills can be gained from repeated patient care experiences. Experiential learning can be attained in a reflective cycle such as Kolb's learning cycle. This cycle applies to patient experiences by observing and reflecting, formulating abstracts and generalizations, and finally testing the implications of concepts in new situations. Theoretical perspective and empirical knowledge are gained from various disciplines including: psychology, sociology, ethics, management, education, and biological sciences.

Experiential education is a methodology where educators/mentors purposefully engage with learners in direct experience and reflection in order to increase knowledge, develop skills, and clarify values. The practical application of experiential learning has to take into account the learner's previous experience, knowledge base, and theoretical preparation. Context of learning will vary, but will guide the learner from being dependent to becoming a supervised participant and then, onto supported independence.

Situated learning is a model of learning in a community of practice where learning is derived from active or guided participation. Guided participation involves cognitive (thinking) ability, problem solving, adequate knowledge, personal skills, and appropriate attitudes and values. Students learn best when involved in genuine professional practice, supported by an experienced, capable mentor. The experienced practitioner (mentor) is the most valuable resource available to the learner. Effective mentor–learner relationships are dynamic and complex and can be achieved through guided participation.

Educational courses on central venous device placement should include basic anatomy, ultrasound physics and imaging, detection/management of major and minor complications, and maintenance/care of the CVAD. Educational programmes reduce the occurrence of CVAD infections and insertion complications. The use of ultrasound, checklists, bundle programmes, and simulation labs in conjunction with organized educational programmes improves patient safety related to CVAD insertion. Experience and skill are also associated with reduced complication rates related to insertion.

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Educational content should be directed at how to perform a CVAD insertion. Clinician knowledge of certain subject areas and competence in device insertion influence the rate of occurrence for CVAD-related complications. This panel of experts recommends the following topics be covered during the didactic section of an educational course on CVADs:

1. Anatomy and physiology of relevant body systems
2. Ultrasound for insertion and assessment
3. Central venous device tip location
(4) Infection control and sterile technique
(5) Device selection and indications
(6) Insertion procedures, complication prevention, evaluation, and management
(7) Care and maintenance practices along with needleless connectors and securement devices
(8) Qualification and competency
(9) Simulation training
(10) Anatomical models
(11) Objective grading and proficiency
(12) Examination and competency
(13) Supervised instruction
(14) Didactic or web-based training
(15) Developing clinical competence
(16) Education for children and neonates

Anatomy
Knowledge of anatomy affects decision-making and may help to avoid insertion complications (inadvertent nerve, arterial, or lung puncture). Inexperience leads to more attempts, more trauma during insertion procedures, increasing the risk of infections. This supports the idea that more training and knowledge of relevant anatomy and physiology reduces complications during CVAD insertions.

In terms of insertion site selection: the vessels, skin, and nerves systems directly impact the ability to insert and maintain a CVAD. Additional body systems include musculoskeletal, cardiac, and respiratory systems. Infection and thrombosis can both be impacted by site selection, skin integrity, and catheter–vein ratio (CVR).

Anatomy and physiology training content:
- Vessel anatomy, location, size, and path
- Vessel differentiation with ultrasound
- Blood flow dynamics
- Virchow’s triad
- Skin integrity, colonization
- Peripheral nerve identification and distribution
- Respiratory anatomy
- Upper and lower extremity, axillary, neck, and chest anatomy

Ultrasound vein assessment and access
Ultrasound-guided access should focus not only on vein puncture but also on assessment and selection of veins based on size, patency, and risk reduction (Table 2).

The use of ultrasound during CVAD insertions increases first-pass success rates and reduces overall complications. Anatomy is variable and therefore, blind insertion or landmark methods have limitations. Ultrasound visualization identifies variant anatomy before the procedure and offers the clinician valuable and potentially lifesaving information. A high-risk procedure can be identified and avoided, and one that would be difficult and traumatic becomes safer for the patient.

It is known that vessel characteristics like size and patency impact vascular access outcomes. A simulation study demonstrated that a CVR that approaches 1:3 reduces blood flow by 80%, while a CVR of 1:4 may only decrease blood flow by 40–60%. At this time, there are no in vivo studies that provide outcomes of CVR. However, the use of ultrasound assessment facilitates vein selection, diameter measurements, appropriate catheter selection, and subsequently reduces the risk of venous thrombosis.

Tip location
Tip location defines a device as central, peripheral, or non-central. A non-central tip location of a CVAD greatly increases the risk for catheter-related complications. Complications include thrombosis, vascular erosion, cardiac erosion, tamponade, and arrhythmia. Even within the superior vena cava, the risk associated with catheter-related thrombosis varies greatly depending on how close the tip is to the right atrium. The more cephalad the CVAD tip is from the right atrium, the greater the risk for catheter-related thrombosis. Defining the exact location of the true cavo-atrial junction (CAJ) has been a subject of much debate. Even varying expert opinions exist on whether a tip should be above or below the CAJ. The tip at or near the CAJ provides the least risk. Some exception for right atrial tip placement may exist for the optimal performance of haemodialysis catheters.

Infection control
Central catheter-related infection remains a high source of morbidity and mortality in the acute and long-term care environment. Through the use of systematic educational
Programmes, compliance tracking, and checklists, strides have been made to reduce the occurrence of infections.\(^8\) \(^{32}\) \(^{40}\)

Infection control and sterile technique are key principles to include in any CVAD insertion programme. The most serious and costly complication of CVAD insertion and maintenance is the catheter-related bloodstream infection (CRBSI). CRBSI has three main sources of contamination: the catheter–skin junction (extraluminal), the needleless connector and hub (intraluminal), and iatrogenic contamination from another site of infection.\(^8\) Understanding the mechanisms of CRBSIs is key to making clinical decisions that mitigate risk including: optimal site selection, the use of maximal barrier precautions, hand hygiene, appropriate use of skin antiseptic solutions, and evaluation of device necessity with prompt removal. These five points are the basic elements of the Central Line Bundle Program proven to reduce the risk of CRBSIs.\(^8\) \(^{32}\) \(^{40}\) The bundle is focused on site selection and the ability of the inserter to maintain a sterile environment during CVAD placement.

Inadequate care and maintenance practices during dressing changes, flushing protocols, access for infusions, and needleless connector disinfection all contribute to the development of complications.\(^8\) \(^{36}\) \(^{47}\) \(^{48}\)

**Device selection and indications**

Device characteristics and indications for use affect the selection of a CVAD and contribute to successful completion of the treatment plan. The size of a catheter influences flow which impacts the risk of vessel wall damage.\(^15\) \(^{29}\) \(^{42}\) There is a debate over whether valved catheters impact device function and decrease overall risk to the patient. Multilumen catheters are considered higher risk for infection-related complications compared with those with fewer lumens.\(^8\) It is incumbent on any education programme to provide current research information on CVAD design features and how these features influence risk profiles.

**Insertion procedure**

The provider of CVAD insertion education shall provide updated evidence-based education regarding device placement based on the current accepted standards and manufacturer instructions for use. Professional organization guidelines\(^69\) and literature\(^10\) give suggestions for proper device insertion and use.

**Care and maintenance**

The Centers for Disease Control recommend that clinicians receive insertion procedure and maintenance education in an effort to prevent device complications.\(^8\) In a study by Davis,\(^49\) 71% of CVAD bacteraemias occurred after 5 days signifying failure and contamination during maintenance practices. The ability to reduce CVAD infections relies as much on care and maintenance as it does on insertion.

**Qualification and competency**

Only clinicians competent in CVAD insertion and maintenance should perform these procedures.\(^8\) Instructors must also demonstrate continued competency and proficiency in CVAD insertion. Competency extends to periodic assessment and not just an initial evaluation at the time of training.\(^51\) A systematic approach is required to continually measure outcomes and provide feedback relevant to competence as a function of safety. Studies evaluating the relationship between clinical knowledge and experience have concluded that the decline in knowledge after initial training is accompanied by a decrease in quality of care.\(^52\)

**Ultrasound and insertion simulation training**

Ultrasound guidance is recognized as the gold standard for CVAD insertion.\(^7\) \(^{18}\) \(^{53}\) All training labs should include ultrasound skills education. At the end of the learning period (see the Developing clinical competence section), all trainees should be able to detect major vessel abnormalities with the aid of ultrasound.\(^12\)

**Anatomical models**

There are a variety of inanimate models useful for simulating vessel anatomy visualization with ultrasound. The best simulation models should include vessels and also mimic the normal body anatomy with muscles, soft tissues, and bones. For this reason, inanimate animal models such as turkey breasts may be effective for simulation practice with ultrasound.

In the future, virtual simulators may be available for perfect hand–eye coordination of the trainee.\(^53\) –\(^55\)

**Objective grading and proficiency**

The application of the learning curve concept to CVAD placement means the reduction, by 50%, of major complications during the learning process. With the landmark technique, the suggested minimal CVAD placements required to achieve minimal experience was \(>50\) insertions.\(^56\) With ultrasound guidance, it has been suggested\(^11\) \(^{12}\) that CVAD-skilled trainees require less training than novices but have more problems with hand–eye coordination.\(^2\) \(^3\) \(^{57}\) –\(^{60}\) There is no common training guideline on ultrasound training.\(^11\) \(^{61}\) This Consensus Task Force recommends 6–8 h of didactic education, 4 h hands-on training on inanimate models, then 6 h hands-on training on normal human volunteers for detection of normal ultrasound anatomy. This training should be followed by supervised ultrasound cannulations, coaching the trainee during the procedure in order to achieve the required minimal skill competence with the lowest rate of complications.\(^62\)

**Examination and competency**

A training log-book should be used by the trainee as a checklist to verify completion of all the steps of the training process. If the trainee fails to pass a step, he/she has to repeat the performance until able to demonstrate competence. The educational competence should be evaluated with a multiple-choice test of at least 100 questions. The trainee must pass with at least 70% (according to similar
Competence in central venous device insertion is sparse. From the literature, it appears that there are no standard methods of assessing the competence of the practitioner. According to recommendations of the National Institute for Clinical Excellence (NICE), those involved in placing CVADs using ultrasound guidance should undertake appropriate training to achieve competence. Defining a competent practitioner in objective terms is a difficult task. Competence is often determined after a pre-determined number of procedures being performed or after a subjective assessment by a proctor who is deemed to be competent. This approach is often criticized however, as it relies on general agreement to determine the number of procedures necessary to gain competence. Other authors suggest an alternative, more objective method of training programme. This assessment process includes the completion of a written test assessing the practitioner’s cognitive level of the procedure. This written exam should be in conjunction with a visual exam to test the knowledge of normal vs abnormal vessels, and a skills test using simulation to test the practitioner’s ability to perform the procedure to a satisfactory standard. The practitioner would then undertake observed procedures under the direct supervision of an experienced practitioner.

The two main approaches to rating technical performance are global rating scales and checklists. The goal of checklists is to guide behaviour by confirming routine. It has been argued that the global rating scales (Likert type scales) do not address procedural shortcomings. Global rating scales of competence have evolved and a validated 24-item checklist using the Angoff method has been created to assess the competence of residents in CVAD insertion. Because CVAD insertion is sequential and predictable, assessment of the procedure can be performed using a checklist, as each component of the skill can be assessed. Formal evaluation of competence is performed using instruments such as the Ottawa Global Rating Scale (GRS) and the Ottawa Crisis Resource Management (CRS) checklist which both demonstrate effectiveness in multiple domains.

Teaching CVAD insertion with children and neonates
Central venous catheterization in paediatric patients and neonates is considered a challenging procedure, even when ultrasound guidance is used. Several papers demonstrate a significant reduction in major complications when ultrasound guidance is used instead of the landmark technique. Central venous device placement in children and neonates requires specific training in order to achieve competence in central vein cannulation.

Materials used for CVAD placement in paediatric patients must follow specific requirements:

(i) The catheter is selected according to the type of therapy to be administered to the child/neonate.
(ii) Ultrasound probes must be linear with a small footprint and a high frequency (from 10 MHz up).
(iii) Nitinol soft-tipped guidewires are preferred.

It is necessary to have minimal cannulation of 70 central vein punctures per year in order to maintain the skills and the competence in these patients. 91 Ultrasound guidance enhances first-time success and time to cannulation in small infants, but requires supervised training to gain proficiency. It has been demonstrated to take 15 supervised, ultrasound-guided cannulations to reach proficiency with small infants. 91

Minimal training is also required for peripheral vein cannulation in paediatric patients. 92 Nurses and physicians should be trained in paediatric-specific i.v. procedures during their academic/employment training period. Even so, additional factors noted in the literature were better indicators of success, such as difficulty and cooperativeness of the child. 93 In cases of difficult peripheral vein access, ultrasound guidance may not always be useful because simple compression of the probe over the site may cause collapse of superficial veins, necessitating the use of a very lightweight probe for these patients. In this case, near-infrared technology has increased success rates of peripheral vein catheterizations. 94 95

Conclusion

Evidence supporting and directing the education and training of those who insert and maintain CVADs is limited. This study suggests evidence-based recommendations that provide direction for establishing consistency in the development of training programmes and measuring competency through completion of didactic lessons, simulation, examination and supervised practice. These recommendations should be considered minimal level requirements for all inserting clinicians. Each recommendation was considered necessary by the consensus group in establishing uniform training throughout the world, both with adults and children/neonates. This document is designed to serve as a guideline for those creating education and training programmes and for those undertaking the training. More research is necessary to establish stronger recommendations and clearer directives.

Supplementary material

Supplementary material is available at British Journal of Anaesthesia online.

Declaration of interest

None declared.

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