
This article provides an overview of IO vascular access and the ways in which this mode of access may be utilized by and interface with radiology. Winthrop University Hospital provides their institutional protocol for using Arrow EZ-IO access for administration of iodinated contrast agent.


This article provides a brief overview of intraosseous access for radiologists followed by a discussion of the use of IO access devices in the radiology suite, particularly for CT imaging. The protocol established at the authors' institution for use of the EZ-IO system is described which emphasizes obtaining confirmation of proper IO catheter placement by use of imagery prior to full infusion of contrast medium.


Preclinical randomized, controlled trial evaluating the relationships between the anatomical distance of IO epinephrine and measures of resuscitative outcome in an adult swine model of ventricular fibrillation (VF). There were no significant differences between the humeral IO, tibial IO, and IV groups relative to the occurrence of ROSC, 30-minute post-ROSC survival, and time to ROSC. The anatomical distance of IO epinephrine injection from the heart did not affect short-term measures of resuscitative outcome in an adult swine model of VF including the occurrence of ROSC, 30 minute post-ROSC survival, and time to ROSC. Rapidly administered epinephrine, irrespective of route of administration, increased the chance of ROSC and survival to 30 minutes post-ROSC in this study.


This article presents a case study of rapid sequence intubation via intraosseous (IO) access with a review of relevant literature. The authors describe a case of an adult male patient, peri-arrest with cardiogenic shock, cyanosed with un-recordable oxygen saturations and blood pressure. IO access was established in the proximal tibia and rapid sequence induction was performed using fentanyl, ketamine and suxamethonium. After 30 seconds direct laryngoscopy was attempted and intubation was secured on first attempt. The authors concluded that use of IO access for RSI can be useful in cases of difficult vascular access and rapid intubating conditions can be achieved which are comparable to using IV drug delivery.


The abstract describes the interim results of an investigational device exemption study evaluating use of EZ-IO in volunteers for a 48 hour dwell time period. At the time of the report, 39 subjects completed the study with no serious adverse event reports. Subjects were randomized to receive IO insertion in the proximal tibia or proximal humerus insertion sites. Pain has been managed using oral hydrocodone/acetaminophen and/or intravenous/intramuscular ketorolac. This study is sponsored by Teleflex Incorporated.


Randomized, prospective preclinical study that examined the differences in pharmacokinetics and pharmacodynamics of tibial IO (TIO) and IV-delivered vasopressin during cardiac arrest and CPR until ROSC was achieved. No difference was noted for ROSC between TIO and IV delivered vasopressin. Authors concluded the use of IO access could avoid the time delay associated with IV access, and that it is effective for treatment of hypovolemic cardiac arrest and should be first line for rapid vascular access.


This article includes a case study of an adult patient who received an intraosseous (IO) catheter, that may have extravasated, resulting in vascular compromise. The patient was treated with pharmacologic intervention and the status was reversed. A review of the literature on adult IO complications is also described.


This study examined the relationship between body mass index (BMI), the ability to palpate the tibial tuberosity (TT), and soft tissue depth at recommended IO insertion sites in obese patients using ultrasound. Authors concluded in obese adults with a palpable TT or BMI ≤ 43, a 25 mm IO needle is likely adequate at the proximal and distal tibial insertion sites; and at the proximal humerus site a 45 mm is recommended.
A preclinical study evaluating blood transfusion via IO vascular access in anesthetized swine. Results showed pressurized blood transfusion through IO vascular access resulted in acceptable flow rates and did not result in appreciable hemolysis as indicated by free hemoglobin values. This study was sponsored by Teleflex Incorporated.

This abstract describes the results of a healthy volunteer study evaluating use of the EZ-IO TALON in the sternal IO insertion site. IO infusion flowrate was measured and reported for gravity infusion, as well as pressurized infusions at 100, 200, and 300 mmHg. The authors concluded the TALON device may be used safely and successfully in the sternum with excellent infusion flow rates. This study was sponsored by Teleflex Incorporated.

A preclinical study evaluating the immediate effects of power injected contrast media on the medullary space of anesthetized swine. Contrast media (150 mL) was administered at a rate of 5 mL/second. For each limb receiving power injection a control limb was submitted for evaluation. The pathologist was blinded to which limb received power injection. Results showed no histological difference in limbs receiving and not receiving power injection. This study was sponsored by Teleflex Incorporated.

This randomized crossover manikin trial compared the NIO and EZ-IO devices for time to placement and ease of use. For both parameters the NIO performed better.

This letter to the editor describes a prospective, randomized, cross-over cadaveric study that evaluated use of the EZ-IO and NIO devices by novice paramedic device users. Following a brief in-service on use of both devices and practice insertions using a leg-trainer manikin, each participant attempted to establish IO access using each device in a resuscitation simulation with an adult cadaver with CPR in progress. Results showed first attempt success rates of 97.4% with the NIO and 100% with the EZ-IO; and mean time to insertion was 16.8 seconds with the NIO and 42 seconds with the EZ-IO.

Preclinical study that examined the differences in pharmacokinetics and pharmacodynamics of tibial IO (TIO) and IV-delivered epinephrine during cardiac arrest and CPR. There were no significant differences between IV versus TIO epinephrine in achieving ROSC, time to ROSC, and Cmax. In the context of ROSC, epinephrine delivered via TIO route was a clinically relevant alternative to IV administration. The authors concluded that when IV access cannot be immediately obtained in cardiac arrest patients, TIO access should be considered.

This article presents a 5-case series describing use of IO vascular access by anesthesiologists in the perioperative and critical care settings. All insertions were made in the proximal tibia and there were no adverse events reported. The devices cited as being used were the EZ-IO and the Cook Surfast manual needle. A proposed perioperative vascular access algorithm incorporating IO access is presented. The authors address key topics around IO access including use of same drug dosing as IV administered drugs, frequent palpation and monitoring of the insertion site for extravasation, low complication rate and actual risks associated with fat emboli and bone injury, pain and anxiety management in the awake patient and clinician-perceived pain. Administration of blood products, ACLS drugs, Lactated Ringer’s solution and anesthetics are noted without complication. Use of IO aspirate for laboratory testing is noted, however use of the initial aspirate is indicated. Several patients in the case series were reported to find the discomfort of IO insertion preferable to multiple intravenous attempts. The authors concluded: IO lines can be placed quickly and safely in emergency situations or in elective surgical patients with difficult intravenous access; IO access can be useful in a wide variety of clinical settings; and is an important skill for anesthesiologist to learn.
This article describes the strategies used at one hospital (Penn Presbyterian Medical Center) to increase the use of intraosseous catheter to rescue patients in all care settings.

This article in Swedish describes a study evaluating use of aspirate obtained from the IO space for laboratory analysis. The authors note that point-of-care equipment should be used for analysis. Creatinine, morphine and troponin was successfully analyzed; leucocytes and platelets were noted to possibly cause falsely elevated values.

A preclinical study in which 8 anesthetized swine were put into an induced septic shock state to allow troponin I level measurements to be compared from serial venous plasma, arterial plasma and intraosseous aspirate specimens collected hourly. Two milliliters of IO aspirate were wasted before collecting each IO specimen for analysis. The levels of IO troponin I increased during the first 3 hours of shock but then plateaued at a high level while the venous and arterial levels continued to increase. Authors concluded that troponin I can be analyzed in bone marrow aspirates in a shock model and that this information may be useful in medical emergencies where cardiac damage is suspected to be involved.

Preclinical study to determine whether intraosseous pressure (IOP) could be consistently recorded and similarity of IOP to central venous and arterial pressure in a porcine hemorrhagic shock model. IOP tracings were tracked reliably from the proximal humerus, distal femur, and proximal tibia. Baseline IOP ranged from 16-18 mm Hg among the three sites, which was approximately 23% of arterial pressure. This study was sponsored by Vidacare LLC.

Literature review on contemporary practices of intraosseous (IO) vascular access in adult patients.

This article presents an overview of IO access in the critically ill adult patient by way of a literature review.

Case report of a prehospital misplacement of an IO catheter into the intra-articular space of the knee joint when access was attempted in the field. Upon ED arrival IO placement was noted to be high and intra-articular placement was confirmed by xray. A sterile NS lavage was done and patient recovered without complication. Authors note this as a previously unidentified complication of IO placement and advise xray confirmation of affected sites with follow-up of intra-articular placements for the septic arthritis. (Picture of site appears to be an EZ-IO).

A cadaveric study performed by twenty-seven medical students, inexperienced with IO vascular access, that compared use of the EZ-IO for access in the proximal humerus and proximal tibia insertion sites and the FASTR for access in the sternum. First pass insertion success, insertion times, and one minute flow rates using external pressures from 0 to 300 mmHg were evaluated. The authors concluded that both the EZ-IO and FASTR devices may be effective IO devices and are likely suitable for fluid resuscitation using a pressure bag.
Bibliography


Retrospective analysis of IO needle insertions performed in all HEMS missions during the first three years (2009-2011) using the EZ-IO® system. Overall success rate of EZ-IO procedures (N=348) was 99.6%, with a first attempt success rate of 85.9%; and high user satisfaction rate of 93%. IO as access was mostly second line overall but first line in children <7, trauma and cardiac arrest. There was 1 failure and 4 needle insertion problems noted; no serious complications.


This single center, prospective, observational clinical study compared use of intraosseous (IO) access to central venous catheter (CVC) access for inpatient medical emergencies, managed by the medical emergency team (MET), within an urban teaching hospital. CVC access training included percutaneous, landmark-guided CVC placement without ultrasound guidance, using the femoral vein as the primary site. For IO access, the proximal tibia was the primary site and proximal humerus was secondary. Results showed IO access was significantly superior to CVC access with regard to first pass success rates, overall success rates, time to placement, and number of attempts for proper placement. On average more CVC kits were used per patient; complications were greater with CVC. There was one serious complication of tissue necrosis secondary to extravasation in the IO group.


A case study report describing administration of prothrombin complex concentrate (PCC) via IO access to treat bleeding caused by Rivaroxaban oral anticoagulant in a 64 year old male. Proximal humerus IO access was established and 1490 units of PCC were administered at its maximum rate of 10 mL/min without adverse event. The same dose that would be administered IV was given IO. The patient experienced pain with IO infusion despite administration of 10 mg of lidocaine and 3 doses of fentanyl 25 mcg given via IO access. The patient was transferred to the medical intensive care unit and ultrasound guided IV access was established. The authors concluded that Profilnine is well tolerated when administered via IO access; however further studies are needed to evaluate if this is an effective practice.


In a healthy adult volunteer study contrast media was injected through the proximal humerus site and captured under fluoroscopy as it entered the heart. The mean time it took from injection at the insertion site to visualize contrast entry into the superior vena cava and the right atrium was 2.42 seconds. Abstract presented at ACEP 2015. This study was sponsored by Teleflex Incorporated.


A simulation study comparing use of manual (Cook Medical) and mechanical (Arrow EZ-IO) intraosseous (IO) devices to establish IO access in mannequin bones representing infant, pediatric and adult tibias. Twenty-two anesthesiologists with no prior experience with IO devices participated in the study. The outcome measures were success rate, insertion time and operator reported difficulty of use. Results were in favor of the mechanical device for insertion time in each category, and success rate in the adult tibia group; there was no statistical difference in the difficulty of use evaluation.


Abstract describing preliminary results for the first 24 subjects of an EZ-IO study evaluating catheter dwell times for 48 hours. Initial data indicate that IO vascular access can be safely maintained for a period up to 48 hours without risk of osteomyelitis or other serious adverse events. Authors also noted that additional analgesics for IO infusion pain management may be more effective than the current solely administering lidocaine into the IO space. This study was sponsored by Teleflex Incorporated.

Overbay JK, Kon AA. Dermal abrasion experienced as an adverse effect of the EZ-IO. The Journal of Emergency Medicine 2015;http://dx.doi.org/10.1016/j.jemermed.2015.09.003

This article presents a case report of a 7 month old female who received intraosseous vascular access via the EZ-IO in the distal femur that resulted in a dermal abrasion where the needle hub contacted the skin. The wound healed without significant complication however the scar at the IO site persisted at 11 months post the event. The authors recommend that providers use the minimal force necessary when operating the EZ-IO to avoid similar adverse events.
A case study report of a 24-year old female who presented to the emergency department after consuming an over dose amount of verapamil. Central and peripheral venous access were obtained for delivery of vasopressors and intravenous fat emulsion 20% (IFE). IFE was initiated via peripheral IV (PIV) access but access was lost; administration through central access was not possible due to the potential drug interaction. Intraosseous (IO) access was established using the Arrow EZ-IO system in the proximal tibia without complication and IFE administration was resumed. The patient reported some pain with infusion. After half the bolus administration was delivered, the infusion pump alarmed due to inadequate flow. PIV access was obtained and IFE administration was resumed using the newly obtained access route. The authors suggested that the viscosity of the medication may have caused the delivery failure by infusion pump through the IO route and recommend slowing down the bolus rate of infusion for clinicians attempting this route for IFE administration in the future.

A healthy volunteer study evaluating use of the EZ-IO TALON in the sternal IO insertion site. Military trained medics performed all device insertions. IO infusion flowrate was measured and reported for gravity infusion, as well as pressured infusions at 100, 200, and 300 mmHg. The authors concluded the TALON device may be used by military and tactical medicine personnel to safely and successfully establish IO access in the sternum with excellent infusion flow rates. This study was sponsored by Teleflex Incorporated.

This preclinical study evaluated the occurrence of fat intravasation resulting from intraosseous (IO) flush and infusion in anesthetized swine. Intravasated fat was assessed using a lipophilic fluoroprobe (Nile red) and by vascular ultrasound imaging. Fat intravasation was observed during all IO infusion regimens, with subclinical pulmonary fat emboli persisting 24 hours post infusion. It was noted that initial flush was a significant factor in fat intravasation, low levels of intravasation occurred with infusions ≤300 mmHg, fat intravasation and bone marrow shear-strain increased with IO infusion rates, and intravasation was influenced by cannula insertion site.

A prsctirical study evaluating the immediate effects of power injected contrast media on the medullary space of anesthetized swine. Contrast media (150 mL) was administered at a rate of 5 mL/second. For each limb receiving power injection a control limb was submitted for evaluation. The pathologist was blinded to which limb received power injection. Results showed no histological difference in limbs receiving and not receiving power injection. This study was sponsored by Teleflex Incorporated.

A retrospective study evaluating attempts to establish intraosseous vascular access in pediatric patients using a manual device and the EZ-IO, in a tertiary care pediatric emergency department. Results showed 35 patients had IO access attempted using manual and EZ-IO devices. In patients greater than and less than 8kg the EZ-IO had a higher success rate but time to placement was longer. Overall success rate including both devices was 64%. There were 2 complications of transient leg swelling after EZ-IO placement in 2 patients.

The authors described a proof of concept pilot study conducted to determine intraosseous (IO) pressure measures and their relationship to blood pressure obtained using an external blood pressure cuff in ICU patients. The average IO systolic blood pressure, IO diastolic blood pressure, and IO mean were 39.5±12.7 mm Hg, 31.5±7.6 mmHg, and 35.0±8.8 mm Hg respectively. The ratio of IO systolic blood pressure to cuff systolic blood pressure, IO diastolic blood pressure to cuff diastolic blood pressure, and IO mean to cuff mean are 34.5±13.4%, 40.5±22.3%, and 40.1±17.1% respectively. There were no adverse events reported. Investigators concluded that in their convenience sample of severely ill and injured patients, IO pressure was reliably obtained and appeared to be 35% to 40% of blood pressure readings obtained via external blood pressure cuff, and that this method of pressure monitoring may be an appropriate alternative to invasive monitoring option in the future. This study was sponsored by Teleflex Incorporated.

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An overview of IO vascular access with a focus on the proximal humerus IO insertion site.


Preclinical study using a porcine model to determine whether there were differences in intraosseous (IO) and intravenous (IV) antibiotic (cefotaxime and gentamicin) concentrations during septic shock. Both methods of administration yielded comparable concentrations.

Authors concluded in an emergency, IO administration of these antibiotics may be considered in severe infections when venous access is difficult.


This case study describes a neonate who suffered a cardiac arrest, had return of spontaneous circulation (ROSC) and was treated with multiple medications and therapeutic hypothermia. The patient had received three IO needle insertions, one in the left tibia that was removed following swelling with bolus injection; one in the left distal femur that dislodged with movement of the patient's legs; and one in the right proximal tibia. Twenty-four hours after initial IO needle placement the child developed pallor and discoloration and was diagnosed with compartment syndrome to the right lower extremity. Five days post-IO insertion a below the knee amputation was performed. Medications infused via the IO access included epinephrine and norepinephrine infusions.


This article in French is a survey of residents and doctors in France that practice in ED, ICU and anesthesiologists units seeking their opinions and practice habits in regard to IO access. Only 29% had ever used an IO kit; with a correlation between years of experience in practice and use of IO access. 555 had received some IO access training; 90% of untrained doctors believed training was necessary. The powered system was the most utilized (EZ-IO).


Literature search for complications associated with IO access included 5759 patients with overall complication rate of 2.1 %. Two cases involving retained needle fragment discussed; one with a proximal tibial EZ-IO that required surgical removal. Authors concluded IO catheters are reliable tools for fluid and drug delivery to critically ill patients with low complication rates (which can be potentially serious but managed).


A prospective observational study that evaluated use of intraosseous vascular access for delivery of rapid sequence intubation (RSI) drugs. Data was collected between January and May 2012 at a combat hospital in Afghanistan. Thirty-four (34) patients underwent RSI with drug delivery via the IO route. Access was established in the proximal humerus and tibia using the EZ-IO and in the sternum using the FAST-1. All placements were successful on first attempt; first pass intubation success rate was 97%; a Cormack-Lehane (C-L) laryngoscoical grade view of 1 was reported 91%. Authors concluded that IO access is a safe and feasible route for delivery of anesthetic drugs for RSI.


Randomized comparative study of adult pigs infused intraosseously with either: 7.5% hypertonic solution (HTS), 3% HTS or normal 0.9% isotonic saline. The animals were observed daily for infection, necrosis and gait up to 5 days, then necropsy and histological analysis was performed for tissue necrosis. Observations included regular tissue morphology and normal gait scores over the 5 day observation period; and absence of gross tissue necrosis and microscopic ischemia post IO HTS infusion in this swine model. Authors concluded this study confirms the clinical safety of IO HTS infusion and its use as an alternative lifesaving treatment.

Preclinical study using a porcine model to determine whether there were differences in intraosseous (IO) and intravenous (IV) whole blood transfusion relative to hemolysis and transfusion time. IO transfusion does not significantly increase hemolysis (using free hemoglobin as outcome measure) or transfusion time compared with IV transfusion. Authors concluded transfusion of whole blood through an IO device is an effective transfusion method that may be used until other vascular access is obtained.


This article presented a general overview of IO use in pediatrics. The history, techniques, anatomy and physiology, complications and a short discussion of most devices on the market, including the EZ-IO, were discussed.


This letter to the editor describes a single case of a needle breaking off after a proximal tibial insertion of the EZ-IO into a volunteer (one of the letter's authors) during a training session. "Divergent from manufacturer instructions the sterile steel stylet was put back into place to achieve better grip for a manual pull-out. Under steady pull in strict axial alignment and gentle clockwise turn, the needle broke away from the plastic connector". The needle was extracted using combination pliers and there is no evidence of damage to the leg. Authors acknowledge this can be avoided by adherence to manufacturer’s directions for use.


This was a prospective, randomized controlled clinical pilot study comparing the BIG and EZ-IO intraosseous (IO) vascular access devices in 52 adult patients admitted to an emergency department with difficult peripheral venous access. Twenty-six patients were randomized to each device; results were first attempt insertion success BIG 92.3%, EZ-IO 84.6% (P=0.668); procedure time: BIG 2.8 ± 1.2 seconds, EZ-IO 5.2 ± 2.2 seconds (P<0.001), significant; difficulty of use (with visual analogue scale): BIG 8.6 ± 6.4 mm, EZ-IO 25.4 ± 12.6 mm (P<0.001), significant. Authors concluded both EZ-IO and BIG are shown to be reliable and safe methods for insertion of intravascular access in emergency conditions. There were no adverse events or complications reported.


Text article that accompanies video featured in The New England Journal of Medicine on intraosseous access which provides a general overview of IO access and demonstration of IO insertion using the EZ-IO and one manual IO needle set.


A cadaveric study performed by dentistry and medical students evaluating the feasibility of gaining vascular access via the anterior mandible bone.


A preclinical study evaluating the bioavailability of antidotes HI-6 oxime and dicobalt edetate when given via proximal tibia intraosseous (IO) access, established via the EZ-IO, compared to intravenous administration via central access in minipigs. Results showed rapid and similar systemic bioavailability of the antidotes when given by both routes and that IO access is an appropriate access route when IV access is impractical.
A preclinical study comparing intraosseous (IO) and intravenous (IV) administration of Hextend in 27 swine for time of administration and hemodynamics. IO access was established in the proximal humerus using the EZ-IO. Results showed time for administration was not significant; there were no significant differences between IV and IO relative to hemodynamics. The author concluded that the IO route is an effective method of administering Hextend.

Manikin study conducted in Poland with 107 paramedic operators designed to investigate the success rate, time of insertion and perceived difficulty of intraosseous access devices during simulated resuscitation using the EZ-IO, Bone Injection Gun and Jamshidi needles. Results were first attempt success: B.I.G.: 91.59%; EZ-IO: 82.66%; Jamshidi: 47.66%; mean procedure time: B.I.G.: 2.0 min ± 0.7; EZ-IO: 3.1 min ± 0.9; Jamshidi: 4.2 min ± 1.0; and ease of use (1-very easy to 5-very hard): B.I.G.: 1.83; EZ-IO: 2.92; Jamshidi: 4.68.

The objective of this study was to evaluate inclusion of IO access in Korean medical education with a selected group of 50 medical students. Students received 1 hour of didactic lecture and a 1 hour hands on session using the EZ-IO and artificial tibias and were tested. Results showed an insertion success rate of 88%. The authors concluded IO access was adequate for medical education in Korea.

This retrospective study reported IO use over a 7-year period during combat operations in Afghanistan by the UK Defence Medical Services. The EZ-IO and FAST1 IO devices were available for use, IO use data was collected from the front line, during helicopter evacuation and at the combat hospital. A total of 1014 IO devices were inserted into 830 adult patients; various medications infused via IO access are listed. Across all cases there were no serious IO complications and 14 minor complications. The author concluded that in the pre-hospital setting in particular and in severely injured trauma patients, IO access should be considered a primary method of obtaining vascular access.

Case study of 36 year-old in septic shock with co-morbidities of IV drug abuse, endocarditis, tricuspid valve insufficiency and pulmonary embolism. Initially impossible to obtain PIV or CVC access; then unable to give desired fluids through 22 gauge PIV when finally placed. Proximal humerus IO access was established with the EZ-IO 45 mm needle set and the patient was resuscitated with 30 mL/kg fluids and multiple medications given in first hour. Conclusions included that CVCs are not always possible and volume treatment with an IO placed sooner rather than later, especially in children but also in adults, can be lifesaving. IO systems should be extensively available throughout the clinical setting. Article in German.

This abstract describes the results of an observational clinical study that evaluated the use of IO vascular access via the proximal humerus insertion site for administration of contrast media for computed tomography examination. Eight subjects were enrolled into the study, 7 procedures were performed successfully with adequate opacification of the images. One subject experienced extreme pain with the contrast injection, the procedure was terminated and an alternative vascular access route was utilized. There were no serious complications reported. This study was sponsored by Teleflex Incorporated.

A preclinical study comparing the time to onset, time to onset peak, and time to recovery of peripheral intravenous and tibial intraosseous administration of Rocuronium. Study results demonstrated there was no statistical difference front the time of administration to complete neuromuscular blockade between the IO and IV administration of Rocuronium; and the recovery of neuromuscular function was significantly longer after IO administration, however it was not deemed clinically significant. The authors concluded that Rocuronium can effectively be used via the IO route without the need for dose adjustments.
This abstract describes a practice seminar held at the 32nd annual meeting of the Japanese Society of Reanimatology for establishing intraosseous vascular access in simulation using the EZ-IO and using Doppler ultrasound to confirm placement. The authors concluded the EZ-IO system enables immediate vascular access to the central circulation and the Doppler method enables objective recognition of needle misplacement.


In a series of studies using healthy adult volunteers the objective was to add to available data comparing IO marrow/blood (initial 1 mL aspirate), IO blood (subsequent aspirate), and venous and capillary blood to determine if there is a correlation between samples for serum lactate and PT/INR levels. Two point-of-care analysers were used. Conclusions were lactate levels obtained from IO blood appear comparable to lactate levels from venous blood; the PT/INR levels did not correlate. This study was sponsored by Teleflex Incorporated.


Retrospective study of the Israeli Defense Force (IDF) registry from January 1999 through October 2012 to identify all cases in which IO access was attempted. The Bone Injection Gun (B.I.G.) was the device used for IO access. A total 37 attempts were made in 30 patients. First attempt success was 53% with an overall success rate 49% when factoring subsequent attempts. Most frequent cause for failure related to providers skill level, and due to the device design allowing little room for error. This study prompted the IDF to seek an alternative for the B.I.G.

Neuhaus D. Intraosseous Infusion in elective and emergency pediatric anesthesia: when should we use it?. Curr Opin Anaesthesiol 2014;27(3):282-7. DOI: 10.1097/ACO.0000000000000069

General review of IO access, with particular attention to perioperative setting and includes published guidelines of the German Scientific Working Group for Pediatric Anesthesia for use of intraosseous access. The author recommends that for children with known difficult venous access physicians discuss the possibility of IO access preoperatively with the family.

Oesterlie GE, Petersen KK, Knudsen L, Henriksen TB. Crural amputation of a newborn as a consequence of intraosseous needle insertion and calcium infusion. Ped Emerg Care 2014;30(6):413-4

Case study of newborn girl resuscitated with 15 mm EZ-IO catheter placed to her right proximal tibia. Medications given included antibiotics, "fluids" and calcium. Demarcation of the infants skin was noted immediately post-calcium administration; with progression to necrosis. Trans-tibial amputation was performed 1.5 months after initial IO access. Authors concluded calcium extravasation most likely caused the injury but were unable to identify extravasation cause; citing possible needle displacement. Cautionary steps to reduce risk emphasized by authors.


This report describes a study conducted by the Air Force Research Laboratory comparing intraosseous infusion rates between IO sites in a cadaveric model to determine if there is a site that is most effective for volume resuscitation. Using 16 cadavers procured within 72 hours of death, IO access was established in the proximal tibia and proximal humerus using the EZ-IO and in the sternum using the FAST1. Results showed the mean flow rate in the sternum was 1.6 times greater than the humerus and 3.1 times greater than the tibia. An abstract describing this report was presented by oral presentation at the 2014 annual scientific assembly for the Eastern Association for the Surgery of Trauma meeting.

This letter to the editor describes a cadaver study performed by 50 interns who had never performed IO insertion, to determine if there is a learning curve associated with use of the EZ-IO for establishing IO vascular access in the proximal tibia. Following training each intern performed 10 IO insertions and were timed. The results showed a difference between the first and the eighth attempts resulting in a decrease in time to insertion by half. The authors concluded that practice insertions are necessary to become comfortable with the device.

Turkey


In this pre-clinical study, 18 units of blood were transfused into 10 anesthetized swine via intraosseous (IO) access. Venous specimens were collected to evaluate free hemoglobin levels as an indicator of hemolysis. Seventeen transfusions were given via the proximal humerus site and 1 via the proximal tibia, using a pressure bag set to 300 mmHg. Mean transfusion flow rate was 61.6 ± 37.3 mL/min and the mean blood volume transfused was 266 ± 74 mL (n=18). The authors concluded that blood transfusion via IO access resulted in high flow rates and did not result in appreciable hemolysis as indicated by free hemoglobin values. This study was sponsored by Teleflex Incorporated.


This article explores use of IO vascular access in combat and tactical settings through a brief review of the literature describing this practice. A small feasibility study is discussed that evaluated the use of cadavers for training 26 U.S. Air Force Pararescuemen (PJs) on establishing IO access in the humeral head using the EZ-IO, both drill-assisted and manually inserted needles. First attempt placement success with the EZ-IO drill was achieved in 25 of 26 attempts; first attempt placement success using the manual occurred in 19 of 21 attempts. The authors concluded that the humeral head IO site is the most appropriate site within the tactical setting; and that use of a human cadaver model for training is an appropriate model.


A preclinical study comparing the recovery of fibrinogen in a porcine model when fibrinogen concentrate is administered via IV and IO access. The study results suggested intraosseous administration of fibrinogen concentrate results in a recovery of fibrinogen similar to that of intravenous administration.


This prospective study sought to evaluate intraosseous flush practices of emergency physicians. Using cadavers, 15 emergency physicians were asked to flush an IO catheter placed in the proximal tibia and proximal humerus IO insertion sites with 10 mL normal saline as they would in clinical practice; IO pressure measurements were recorded using an IO catheter inserted in the diaphysis of the target bones. Results showed the median IO pressure generated was 903 mmHg and the median flush duration was 5.2 seconds. Result showed significant interoperator variability with greater than 35-fold difference in flush forces. The authors concluded that it may be prudent practice for providers to extend the flush over several seconds to limit the maximal pressures.

Vassallo J, Horne S, Smith JE. Intraosseous access in the military operational setting. J Royal Naval Medical Service 2014;100.1:36-9

This article describes a prospective, observational study conducted March – July 2011 at the emergency department, Camp Bastion, Afghanistan evaluating use of IO access in 117 patients established using the EZ-IO and the FAST1 devices (76% EZ-IO).


Abstract presented at the Society of Cardiovascular Computed Tomography on preliminary findings of an observational study done after training ER physicians and techns on intraosseous (IO) catheter use and implementation of a policy for IO access use. Authors report high injection rates and excellent computed tomography angiography (CTA) scans safely with use of an IO for power injection of iodinated contrast media (ICM). Authors concluded cardiovascular imaging physicians, surgeons, ER physicians, and CT technologists should be familiar with the techniques of IO needle placement and use for power injection of ICM for CTA. The diagnosis and treatment of critically ill and unstable patients may be hastened by this technique.

Case report of 54-year-old male obtunded patient requiring a CT angiogram to diagnosis a suspected massive pulmonary embolism. After several failed attempts to reestablish PIV access, 150mL of contrast were injected through the proximal tibia IO catheter placed by EMS. Excellent opacification of the pulmonary arteries was achieved and there were no immediate complications from the injection noted.


In a letter to the editor this study reports data collected (during a survey of one third of academic emergency medicine programs in the U.S.) regarding IO use in adults and comparing IO access with other vascular access techniques through simulation. Data suggest that IOs were used less than 5% of the time patients needed emergent access and a peripheral line was unobtainable. The EZ-IO was most often used IO device. Authors conclude IO use should be considered more frequently in critical, unstable patients. (This research was presented at the ACEP Research Forum in 2010).


This letter to the editor describes a prospective, observational, trial that evaluated use of the EZ-IO in critically ill and injured patients (adult and pediatric) in a multijurisdictional prehospital setting; 9 EMS agencies were included. The 25mm needle set was the only needle size allowed for the study. One-hundred-eleven prehospital IO placements were performed by EMT-intermediates and EMT-Paramedics with 96 successful placements (86.5%); the most common cause for failure reported by the author was thought to be patient obesity and inadequate needle length. Cardiac arrest patients made up 74.7% of the study population and the most common site accessed was the proximal tibia. Device operators rated the ease of use 7.87 using a 0 to 10 scale where 10=extremely easy.


An observational clinical study evaluating use of the EZ-IO in patients requiring urgent vascular access that would have otherwise received a central venous catheter due to a lack of other options. One hundred five (105) patients were enrolled across five hospitals. The authors concluded that use of IO access in place of CVCs provides time savings, safety, ease of use, and is effective at significant cost savings; IO access may be used as a bridge to CVC placement under optimal conditions; and IO access may be used to replace use of CVCs all together in selective patients. This study was sponsored by Vidacare Corporation.


This article describes a mannequin and cadaver study that evaluated use of the EZ-IO sternal device and the Illinois needle to establish sternal IO vascular access by dental students. Results of the cadaver study showed two cases of perforation of the posterior sternal cortex when the Illinois needle was used and one EZ-IO insertion in the soft tissue without entering the IO space. The authors concluded use of the EZ-IO sternal device with the insertion site template and scalpel incision may be more efficient and less predisposed to complication than use of the Illinois needle.


A letter to the editor reporting a case study of skin necrosis after IO administration of norepinephrine following resuscitation of a 74 years old in septic shock. The EZ-IO was placed to the proximal tibia; approximately 45 minutes post- norepinephrine administration symptoms of necrosis were evident. Authors cite 3 hypotheses for the cause of necrosis and consider that amines’ high level concentration could induce local toxicity in the bone matrix and artery spasm; suggesting it is necessary to define an upper limit of amines’ concentration that should be administered through IO vascular access.


A pre-clinical study that evaluated use of intraosseous (IO) pressure as an indicator of changes in fluid volume status during a hemorrhagic shock protocol. Central venous and arterial pressures were used as comparators. Results showed IO pressure decreased consistently during the controlled shock protocol. Authors concluded IO pressure appears to be equivalent to CVP as an indicator of fluid volume status. This study was sponsored by Vidacare Corporation.
Bibliography

A pre-clinical study that compared intraosseous (IO), central venous and arterial pressure tracings in a porcine model. Results showed that IO pressure was approximately 25% of arterial pressure. A sampling of IO blood gases revealed oxygenation levels of venous blood. This study was sponsored by Vidacare Corporation.

A case study describing intraosseous pressure monitoring, through tibial IO access, using a standard arterial pressure monitoring transducer during resuscitation of a 31-year-old male in cardiac arrest. Pressure readings were recorded for approximately 53 minutes and were compared to non-invasive blood pressure cuff monitoring at the same time points. IO systolic, diastolic and mean IO pressures were approximately 40% of arterial pressures. This is the first case report demonstrating IO space has a measureable blood pressure and it correlates with pressure obtained through conventional techniques.

General discussion on use of the intraosseous vascular access route for infusion of CT contrast, with attention to clinical considerations pertinent to nurses working in the imaging suite. Author also reviews general IO principles and IO devices.

This is a preclinical study comparing the EZ-IO and the Cook manual IO needle when used by 21 resident physicians to establish IO access in anesthetized swine. Results showed the drill-assisted needle was successfully placed 100% of attempts vs 76.2% successful placement with manual; time to placement and user preference also favored the EZ-IO. Technical issues reported included bending of the manual needle 33% of attempts.

A questionnaire study in which Scandinavian emergency physicians, anesthesiologist and pediatricians reported complications they have experienced with IO vascular access based on recollection alone. Complications were reported related to establishing IO access and using established IO access. Out of 1,802 IO cases reported by 386 responders, the most frequently reported complications included difficulty with peristeum penetration and bone marrow aspiration when establishing IO; and slow infusion and needle displacement with established IO access. Osteomyelitis and compartment syndrome were reported with an occurrence of 0.4% and 0.6%. Researchers concluded that potential complications following IO insertion should be addressed during training. Devices discussed included the EZ-IO, BIG, Cook-Surfast, and other unidentified IO devices

This article in German explores use of intraosseous access in a dental practice emergency. In a simulation study, dental students attempted to establish standard peripheral IV access and IO access using 3 different devices: EZ-IO, BIG, and manual IO. Results showed the manual was the fastest to insert, however, the EZ-IO had the highest first-attempt success rate as well as the lowest number of total attempts to IO access.

This abstract describes a study in which 66 obstetric anesthetists, obstetricians and midwives were training on the EZ-IO and evaluated for successful application of the skill in a mannequin study. The participants also completed a survey following their insertion attempt regarding their perceived ease of use and likeliness to consider IO use in the future. Results showed first attempt success was 95.5%; respondents indicated they found the EZ-IO to be easier than establishing PIV access and 100% indicated they would consider IO use in the future.

UK
**Bibliography**


This abstract describes the results of an online survey taken by members of the Obstetric Anaesthetists’ Association, evaluating use of IO access in obstetric emergencies, and availability of IO equipment on UK labor wards. Results showed many members are trained on IO access, consider it a viable option during emergencies however have limited access to equipment.

UK

**Kim S. Intraosseous access: an important clinical procedure for emergency physicians. Lifeline 2013;June:12-3**

Article featured in June 2013 issue of California’s ACEP monthly newsletter. This article discuss general IO principles with examples of several short case reviews and highlights the EZ-I0.


Doi:10.3109/10903127.2012.755582

Pre-clinical study comparing flow rates achieved after insertion with the EZ-I0 in the proximal tibia, distal femur, and proximal humerus in a swine model. IO catheters were placed in each site and normal saline was infused for 10 minutes using a pressure bag at the highest achievable pressures greater than 300mmHg. The flow rates through the proximal humerus were statistically greater than that of the femur or proximal tibia. The femur flow rates were higher than the proximal tibia but similar. Post-mortem histopathologic evaluations done to assess for damage due to the high infusion pressures were consistent with IO catheter placement.


In this pre-clinical study, investigators sought to determine if the pressure readings at the proximal tibia IO site served as a good indicator of proper IO placement when the foot of the limb was squeezed. Traditional methods used to determine correct IO placement, including needle stability, aspiration of blood, and easy infusion, were used as comparators. Results showed the increased pressure reading at the IO site successfully predicted correct IO placement in all cases; traditional methods did not consistently correctly identify proper IO needle placement.


This article provides an overview of various vascular access modalities in emergency medicine including peripheral IV, venous cut-down, central venous catheter, intraosseous access, umbilical vessel access, and arterial access. The anatomy and physiology, indications and contraindications, procedure steps and special considerations are outlined for each access methods discussed.

**Lyon RM, Donald M. Intraosseous access in the prehospital setting-Ideal first-line option or best bailout?. Resuscitation 2013;84:405-406. http://dx.doi.org/10.1016/j.resuscitation.2013.01.027**

Editorial reviewing a case series of EZ-I0 use in the pre-hospital setting in Switzerland by Santos et al., combined with a literature review. The authors conclude IO access should probably be used selectively and training on its use improved, insertion sites should be compared and further investigation of use of the EZ-I0 in major trauma patients, infusing blood components, use in infants, and application of training warrant further investigation.


This abstract describes a study in which the investigators sought to determine the approximate patient population in which the 25mm EZ-I0 needle set was sufficient length to establish IO access in peripartum patients. Ultrasound was used to determine the tissue depth at four insertion sites. Twenty-six women were recruited with a median gestation of 34 weeks. In 88% of patients with a BMI<40 kg/m² the 25mm needle is sufficient to reach the bone marrow at both tibial sites. For the humeral site, IO placement may be more difficult for patients with a BMI>25 kg/m².

UK


A clinical study evaluating the relationship between IO blood and peripheral venous blood lactate levels analyzed using the i-STAT point-of-care analyzer in healthy volunteers. Results showed IO blood lactate levels were comparable to venous blood lactate levels with a positive statistical correlation. This study was sponsored by Vidacare Corporation.
Bibliography

Oksan D, Ayfer K. Powered intraosseous device (EZ-IO) for critically ill patients. Indian Pediatrics 2013;50(7):689-91

A retrospective chart review evaluating use of the EZ-IO in 25 pediatric patients between July 2008 and August 2010 at a Turkish university affiliated hospital. All attempts were made in the proximal tibia and IO access was attempted following failed PIV access within 60 seconds. First attempt success was 80%; the most reported complication was simple extravasation (3 cases) and needle dislodgement (1 case).


This article in French gives an overview of intraosseous vascular access including the physiology of IO infusion, insertion sites, indications, and complications. Available IO devices on the market are described including, time to insertion, success rate and cost.


A case study describing use of the EZ-IO in Afghanistan by US military on 5 patients with traumatic injury including one pediatric patient. Access was obtained in the proximal tibia on first attempt and was used to administer crystalloids in all patients along with opioids, analgesics and antibiotics. All ultimately received central venous access and peripheral access was established in one patient. There were no IO complications.


An observational study evaluating use of the EZ-IO by two ground and one air based physician staffed EMS and at a German surgical university hospital between January 1, 2008 and December 31, 2011. The EZ-IO was used to establish IO access 88 times in 87 patients; 83 insertions were performed in the EMS and 5 were performed in the hospital. The proximal tibia was the primary site used (97.7%) and the first attempt success rate was 94%. IO access was the first approach for vascular access in children compared to adults (38.9% vs. 86.2%). There were 5 failures attributed to missed insertions or extravasation and 2 for wrong needle length. There were no serious complications.


This article reviews the clinical effects of both high-quality chest compressions and the effects that interruptions during chest compressions have clinically on patient outcomes. The authors indicate intraosseous vascular access should be heavily considered as the first or at least second-line method used to help prevent prolonged compression interruptions for the purpose of establishing vascular access. The authors note that when using the EZ-IO this method of access is fast, effective, can handle all resuscitation fluids, and can minimize no flow time when used properly.


A quality initiative study conducted evaluating use of the EZ-IO needles in pediatric patients and their associated complications rates when placed by EMS/ED staff compared Air Evac Lifeteam placement in 2012. The authors concluded that the powered IO device was appropriately utilized by ED/EMS staff as well as Air Evac staff and that there was no difference in the complication rate when the device was used by the two groups.


This article describes a case study of a 5-month old infant that suffered a head injury resulting in shock. She received 100 mL of red blood cells via the EZ-IO in the proximal tibia, resulting in rapid hemodynamic improvement. A literature search was completed for cases of IO blood transfusion in pediatric trauma. Authors note IO availability and knowledge play an important role in hemorrhagic shock; and RBC infusions via the IO route are feasible in this age group.


Article in German

This randomized, controlled study compared tissue concentrations at the surgical site of regionally and systemically administered prophylactic vancomycin, in 30 patients undergoing total knee arthroscopy. The antibiotic was administered using three methods: 250mg through IO regional administration in the proximal tibia (IORA); 500mg through IORA; and 1g administered systemically through IV. Results showed the tissue concentration of vancomycin was greater in the 250mg IORA group than the systemic IV group, and the 500mg IORA group had higher concentrations than both groups.

YEAR: 2012


A pre-clinical study that compared the EZ-IO 15 gauge 25mm needle set and the 13 gauge Jamshidi aspiration/biopsy needle when used to obtain core biopsy specimens in canines.

Canada


This article describes a questionnaire study that was given to UK Role One military clinicians deployed to Afghanistan to assess the level of experience and confidence rating with use of IO access, using the FAST-1 and EZ-IO, and IV access. Thirty-three responses were received; clinicians felt more confident with IV access over IO access; clinicians felt more confident with FAST-1 IO access than EZ-IO IO access.

UK


Preclinical study using a porcine model comparing the maximum concentration and time to maximum concentration of epinephrine administered via the tibial IO, sternal IO and IV routes during CPR. The IV route of administration of 1mg of epinephrine resulted in a serum concentration 5.87 and 2.86 times greater than the tibial route and sternal route respectively. The times to peak concentration was similar for IV and sternal IO groups but delayed for the tibial route. Authors conclude that due to limitations of their study the guidelines of administering 1mg of epinephrine via the IO route should not be changed; further studies using larger sample size, larger volume flush, arterial blood samples and the use of a more precise method of measuring serum epinephrine should be done.


This article discusses how IO access can be used to improve advanced life support therapy. The EZ-IO is described in this article; published comparative studies between other IO devices and peripheral IV access are cited, leading the author to conclude the EZ-IO is user friendly, and establishes intravascular access more quickly and more often on first attempt than other devices.


This pre-clinical study evaluated IO flow rates obtainable with infusion of lactated Ringer’s and hetastarch 6% through the proximal tibia and sternum IO insertion sites, using a swine model. The EZ-IO 25mm was used to facilitate tibial IO access; sternal access was established using a Jamshidi needle. Results showed that hetastarch flow rates were lower than lactated Ringer’s flow rates at both insertion sites; sternal infusion of hetastarch is likely to provide greater estimated intravascular volume expansion than lactated Ringer’s, despite the lower infusion rates; resuscitation using the IO rate is likely to benefit from pressure bag or high-pressure pump delivery. This study was sponsored by Vidacare Corporation.


This abstract presented at the 2nd World Congress on Vascular Access 2012 reports data collected on the knowledge gaps and barriers to IO vascular access use. Two focus group discussions were held at professional conferences (American College of Emergency Physicians and the Emergency Nurses Association) and facilitated by clinical researchers. Data was qualitatively evaluated and researchers identified several main areas of concern for clinicians in both implementation and knowledge gap areas. This study was sponsored by Vidacare Corporation.
Coutry L, Hssain I, Joshi G, Diemunsch P. Intraosseous access for fluid administration in a simulation setting: Comparison with intravenous access. American Society of Anesthesiologist Annual Meeting 2012; abstract number A895. This simulation study compared intraosseous (IO) vascular access, via EZ-IO, with peripheral venous (PIV) access for time to access, perceived ease of placement, rapidity, and safety, and which will be first choice in life threatening situation among 73 prehospital care providers with no prior experience with IO access. Results showed time to placement for IO access was significantly faster than that of PIV; the majority of device operators graded the device superior to PIV for ease of placement, rapidity and safety.

Duncan L, Cohen J, Triner W, Rea J, King C. Intraosseous administration of CT Contrast in a porcine model: a feasibility study. Ann Emerg Med 2012;60(4S):S92 This abstract presented at the 2012 ACEP Research Forum discusses a swine pre-clinical study evaluating CT image opacification when contrast is delivered via IV and proximal humerus IO access. The EZ-IO was used to facilitate IO access. Results showed that under blinded radiology review the IV and IO images were judged adequately opacified to meet diagnostic criteria. Authors concluded that IO administration of contrast material may be a viable alternative if other vascular access is unavailable or if establishing other access will lead to a delay in diagnostic evaluation. This study was sponsored by Vidacare Corporation.

Esteo OV. Intraosseous access in prehospital emergency care. Emergencias 2012;24:44-6 A prospective, observational study which evaluated use of the EZ-IO within the prehospital setting over the course of a 3 year period, in Barcelona, Spain. Included patients were in cardiac arrest or with hemodynamic instability, without peripheral venous access after 90 seconds or 3 attempts to establish access. Results showed IO access was attempted in 49 pediatric and adult patients with an overall success rate of 93.9%; complications included extravasation and pain. IO access sites included the proximal tibia (71.4%), proximal humerus (22.4%) and distal tibia (6.1%). The author concluded that IO access is a viable access route for the management of critical patients or those in cardiac arrest in the pre-hospital setting due to the ability to obtain rapid access and the high first attempt success rate.

Hoskins SL, Nascimento P Jr., Lima RM, Espana-Tenorio, JM, Kramer GC. Pharmacokinetics of intraosseous and central venous drug delivery during cardiopulmonary resuscitation. Resuscitation 2012;83(1):40-5. doi: 10.1016/j.resuscitation.2011.07.041 Pharmacokinetics of IO drug delivery was compared using the tibia or sternum, versus central venous delivery during CPR. Anesthetized swine with KCl arrest were used for this study, CPR was initiated 8 minutes post arrest. Using 2 study groups, dye was injected as a bolus with adrenaline through either the IO sternal and tibial needles or through the IO sternal and IV central venous needles. Results showed peak arterial blood concentrations were faster for sternal IO vs tibial IO administration. Tibial IO delivered 65% of the total dose delivered with sternal administration. Peak blood concentrations were similar for sternal IO and central venous administration. Sternal IO delivered 86% of the total dose delivered by central venous administration. The EZ-IO and Jamshidi were used to facilitate IO access. This research was sponsored by Vidacare Corporation.

Ibrahim M, Cairney K. Intraosseous (IO) infusion as a means of vascular access. British J of Resuscitation 2012;Autumn:23-6 This article provides an overview of intraosseous vascular access, including applicable patient population, IO access sites, pain management, IO education and compares IO access to central venous access.

Kalechstein S, Permual A, Cameron BM, et al. Evaluation of a new pediatric intraosseous needle insertion device for low-resource setting. Journal of Pediatric Surgery 2012;47:974-979. doi: 10.1016/j.pedsurg.2012.01.055 This article describes a study evaluating a new manual needle insertion device, the Near Needle Holder, which uses hollow-bore needles to establish IO access. In a comparative study, healthcare professionals attempted IO insertion in the proximal tibia insertion site of a mannequin using the NNH and a standard Cook manual IO needle. Participants then completed a questionnaire regarding their experience. The most reported complication was the plunging of the needle into the medullary space from the decrease in resistance once the cortex was penetrated. Other IO devices on the market are mentioned, including the EZ-IO.

Landy C, Plancade D, Gagnon N, Schaeffer E, Nadaud J, Favier JC. Complication of intraosseous administration of systemic fibrinolysis for a massive pulmonary embolism with cardiac arrest. Resuscitation 2012;83:e149-50. doi: 10.1016/j.resuscitation.2012.01.044 This letter to the editor describes a case in which a 53-year-old male in ventricular fibrillation received IO access via the EZ-IO in the ED with suspected massive pulmonary embolism. The patient was successfully resuscitated. Necrosis of the anteromedial side of the leg, at the IO site, presented 48 hrs post IO use. After 18 weeks the patient underwent surgical grafting. The authors linked the necrosis to adrenaline extravasation and local ischaemia. While the authors conclude that thrombolysis or repeated high doses of adrenaline should be given via the IO route when needed, it is not without the risk of complication.

This article provides an overview of intraosseous vascular access, including applicable patient population, IO access sites, pain management, IO education and compares IO access to central venous access.

This clinical trial evaluated the time required to establish IO access versus central venous catheter (CVC) in adults undergoing resuscitation, who had failed peripheral IV access (PIV) attempts. IO and CVC placement were performed simultaneously; two IO devices, the EZ-IO and the BIG, were used to facilitate IO access in randomized format. Forty (40) patients were enrolled, first attempt success for IO was 85% vs 60% for CVC placement; median procedure time was 2 minutes for IO vs 8 minutes for CVC. The author concluded that though IO access is safe, reliable and rapid during resuscitation, it cannot replace CVC but should be considered as a valuable bridging technique.


This abstract presented at the 2012 NAEMSP scientific assembly described a randomized, cross-over study in which 8 swine were administered chilled saline via IV and IO routes to determine if the two routes were equivalent. The results suggested no clinical or statistical difference between IV and IO routes for infusion of chilled saline for therapeutic hypothermia. This study was sponsored by Vidacare Corporation.


This abstract presented at the 2012 ACEP Research Forum discusses a literature review of intraosseous access publications since 1985 providing an updated safety profile for IO access. The search resulted in 192 articles describing IO access with 6 cases of osteomyelitis and 6 cases of compartment syndrome secondary to extravasation reported. Of the 192 articles identified, 140 described the EZ-IO. This study was sponsored by Vidacare Corporation.


This abstract presented at the 2012 ACEP Research Forum describes a preclinical swine study evaluating the ability to induce therapeutic hypothermia by infusing chilled saline via IV and IO access. The EZ-IO was used to facilitate IO access. Results showed statistical equivalence between IV and IO routes when used to deliver chilled saline to induce therapeutic hypothermia. Results also showed that use of chilled saline and infusion tubing submerged in an ice water bath provides the most effective means of cooling. This study was sponsored by Vidacare Corporation.


This abstract presented at 2012 NAEMSP scientific assembly described a retrospective study that evaluated success rates and features of prehospital IO placement in adults following implementation of the EZ-IO, over a 2 year period. First attempt success rate in 281 patients was 89.7%; overall placement success was achieved for 91.8%.


A literature review of articles describing intraosseous vascular access devices used in the pre-hospital setting. Twenty articles met the inclusion criteria and described the manual devices, BIG, Fast-1 and the EZ-IO. The authors concluded that the literature suggests that semiautomatic IO devices may be more effective than manual devices.

Page D. Intraosseous intrigue: Studies examine success rates on pediatric, adult & obese patients. JEMS January 2012;32-3

In this article, the author discussed five recent studies on intraosseous access providing his opinion about the quality of each study.


An overview of IO vascular access including a review of currently available literature. The author discusses various IO devices available and their performance metrics, IO access sites, flow rates, advantages and disadvantages of IO access compared to conventional access methods, complications and recommendations on use of the approach. The author concludes that while IO access may not be appropriate for all patients, it deserves a place in the modern provider’s armamentarium.

This letter to the editor describes a case in which sternal IO access was established using a Jamshidi needle to administer iodinated contrast for a thoraco abdominal CT on a 61-year old male who presented to the ED with respiratory distress. Picture quality was deemed excellent by the radiologists. The authors conclude that the sternal IO route can be used with excellent picture quality but it should be used only in exceptional cases due to the potential risks of a high-power injection through the bone. EZ-IO is mentioned as an alternative IO device available.


An article discussing the technique and safety profile of intraosseous access using the EZ-IO device. Needle selection, contraindications, insertion sites and techniques, catheter stabilization and removal are all discussed along with the safety profile of the EZ-IO against other IO devices and central venous catheters. The authors concluded that IO access should be considered whenever immediate vascular access is required. This article was co-written by an employee of Vidacare Corporation.


This abstract presented at the 2nd World Congress on Vascular Access 2012 describes the results of an analysis of published IO complications since 1985. The safety profile of the EZ-IO is also discussed in this abstract. The authors conclude that new devices and techniques have resulted in an approved IO safety profile. This study was sponsored by Vidacare Corporation.


This article presents a general overview of intraosseous (IO) vascular access in the pediatric population through an IO literature review. Available IO devices were discussed.


This pre-clinical study sought to evaluate the various pressure levels obtained by 22 veterinary clinicians when administering a 10ml normal saline flush of an IO catheter. The EZ-IO was used to establish access in an isolated, cadaveric swine femur. The authors found the median peak intraosseous pressure was 615 mmHg with a range of 57 to 1,100 mmHg. Authors concluded that there is a great deal of variability between clinicians and their flush pressure and that a standardized flush protocol may be beneficial.


This article describes a retrospective study in which 50 consecutive MRI images were evaluated of the humerus for the purpose of determining the optimal needle length necessary for successful proximal humerus IO insertion. Results showed the cortical thickness was 4mm in all cases and that an IO needle length ranging between 40-50mm should be used via the anterior approach. The EZ-IO is specifically discussed in relation to the proximal humerus IO insertion site; and a 24 patient post mortem review of the EZ-IO placed in the proximal humerus is discussed.


An observational study evaluating use of the EZ-IO in a Swiss pre-hospital EMS system between January 1, 2009 and December 31, 2011 and comparing those results to the literature. Sixty IO insertions were performed on 58 patients; the proximal tibia was used in all attempts except 1 attempt made in the proximal humerus. Overall success rate was 90%; the 6 failures were attributed to impossibility to infuse, difficult needle insertion, and incorrect insertion site (tibial plateau). Some of the indications for IO access included cardiorespiratory arrest, major trauma, and shock; general anesthesia was successfully inducted in 7 patients. Drugs infused are listed. There were no serious complications.


This prospective observational study compared flow rates between distal and proximal tibia IO access in adults, with each adult serving as their own control. The EZ-IO was used to facilitate IO access. IO infusion was performed with and without pressure. The authors concluded that infusion flow rates were significantly higher in the proximal tibia as compared to the distal tibia, and that flow rates are significantly higher with pressured infusion vs. non-pressured infusion. This study was sponsored by Vidacare Corporation.
Torres F, Galán MD, Alonso MD, Suarez R, Camacho C, Almagro V. Intraosseous access EZ-IO in a prehospital emergency service. Journal of Emergency Nursing 2012;http://dx.doi.org/10.1016/j.jen.2012.03.005
This observational pre-hospital study conducted in Madrid, Spain prospectively evaluated use of the EZ-IO Jan 2007- Dec 2009 as an alternative to peripheral IV access. During the study period, 107 patients underwent 114 EZ-IO insertions and all were successful on first attempt. IO access was established in the proximal tibia (49%), distal tibia (25.2%), radius (14.9%), and humerus (10.5%) and all lines were the first form of vascular access established in the patient. There were no adverse events or complications.

This abstract presented at the 2012 ACEP Research Forum explored the viability of the distal femur as an IO insertion site with a literature review of IO vascular access and brief overview of a post-mortem study of pediatric distal femur insertion success. Authors concluded that clinical literature, clinical studies, and a post-mortem study confirm that the distal femur is a viable option for IO infusion in pediatric patients. This study was sponsored by Teleflex Incorporated.

This poster presented at the 2012 International Conference of Emergency Medicine described a 4 month review of intraosseous access in UK military operations in Afghanistan. During the timeframe the EZ-IO was used to establish IO access in the proximal humerus and tibia sites; the FAST1 was used to establish sternal IO access. Of the 87 EZ-IO applications there were 12 functional issues and the placement success rate for both sites combined was 86.3%. In 24 FAST1 applications there were 4 functional issues and the placement success rate was 83.4%.

In this article the authors review the evidence supporting the use of IO access; determine the utilization IO access as described in the literature; and assess the level of specialty society support. Various IO devices are mentioned including the EZ-IO.

doi:10.1016/j.ajem.2011.07.010
This study conducted by the San Antonio Fire Department evaluated the first-attempt success rate for humeral EZ-IO placement by paramedics in prehospital adult cardiac arrest patients. Humeral placement was attempted in 247 cardiac arrest patients; first attempt placement success rate was 91%. Authors concluded that humeral IO placement is a reliable method for vascular access in this patient population. This research was sponsored by VidaCare Corporation.

A literary search of electronic databases was performed to identify publications comparing IO access devices. Publications qualifying for study evaluation must have compared two or more semi-automatic IO devices or at least one semi-automatic device and a manual device. Reviews, editorials, surveys, and case reports were excluded. Ten comparative studies met the qualifications for inclusion and are briefly discussed. The studies evaluated suggested superiority of the batter powered IO driver over manual needles and other semi-automatic IO infusion devices.

A clinical study comparing Cefazolin concentrations found at the operation site following total knee arthroscopy when prophylactic antibiotics are administered systemically, through IV administration, and regionally, through IO injection directly at the site using the EZ-IO. Subcutaneous fat and bone samples were collected for evaluation from 22 subjects. Authors concluded that regional IO administration of prophylactic antibiotics can achieve tissues levels higher than with systemic administration.

YEAR: 2011

Auerhammer J. Lebensbedrohliche arterielle blutung aus der a. carotis communis: Fallstricke bei der intraossaren punktion. Notfall Rettungsmedizin 2011;14(2)147-150;doi 10.1007/s10049-010-1380-1
This article in German presents a case of a 67-year-old female patient with an arterial bleed and venous access difficulties in whom IO access was attempted unsuccessfully two times using two different IO systems. The author concluded that IO success is dependent upon IO anatomy and physiology knowledge as well as knowledge of the device being used.

9/6/2016

This article describes an animal trial that assessed the ability of protected, experienced first responders and limited-experience first receivers to place IO lines for antidote administration using the Vidacare EZ-Io device. First responders placed IO lines successfully in 100% of cases, and first receivers placed IO lines successfully in 91% of the cases. Investigators concluded that IO lines may facilitate earlier administration of antidotes to hazardous material victims.


This article in Spanish describes an IO complication case in which a newborn infant developed tissue necrosis as a result of extravasation during IO infusion.


The case report describes a woman experiencing massive hemorrhaging following emergency caesarean delivery. Though the patient possessed a peripheral IV catheter, additional IV access was needed and gained through the proximal humerus IO space using an EZ-Io. This vascular stabilization and additional filling of the central volume through the IO route allowed placement of a subclavian central line. Authors concluded that a key to the resuscitation process was the rapid utilization of the IO.


A case study report in French describing compartment syndrome secondary to intraosseous infusion in a 57-year-old burn patient. IO access was established in the proximal tibia on second attempt; both attempts were made in the same limb though it was noted that the first attempt did not penetrate the cortex. Drug and fluid infusion was initiated; ten hours later the limb was found to appear ischemic. The IO catheter was removed and compartment release was performed. The author concluded that IO access remains an important mode of vascular access and that adherence to contraindications and careful clinical monitoring should decrease risk of complications.


This article presented a general overview of IO use in pediatrics. The history, techniques, anatomy and physiology, complications and a short discussion of most devices on the market, including the EZ-Io, were discussed.


An overview of available intraosseous vascular access devices, including the EZ-Io.


Case study of a 42 year-old woman with massive obstetric hemorrhage ultimately resulting in postpartum hysterectomy. Massive blood loss and inability to stop bleed prevented sufficient resuscitation via established PIV lines. IO access was established with the EZ-Io and used for fluid replacement and administration of cardiac resuscitation drugs. Fluid administered through IO access was 75% of the total infusion volume.

Dolister M, Miller ST, Borron S, Truemper E, Shah MR. Intraosseous vascular access can be used safely and effectively, and at a lower cost than central venous catheters, for pediatric and adult patients in the hospital setting. Ann Emerg Med 2011;58(4S):S311

This abstract describes the interim results of an observational clinical trial evaluating use of the EZ-Io to establish venous access in patients that would typically receive a central line due to lack of other options. At interim analysis, 50 patients had been enrolled in the study. First attempt IO access success rate was 96%; mean time to IO access was 95.1 seconds. The authors concluded that IO access in place of or as a bridge to central venous catheters is safe, fast, and effective for adult and pediatric patients in the hospital setting with substantial cost savings potential. This research was sponsored by Vidacare Corporation.
German Society of Anaesthesiology and Intensive Medicine eV (DGAI), includes a general discussion of intraosseous (IO) vascular access, overview of devices and recommendations for pediatric anesthesia with indications for intraosseous infusion in pediatric anesthesia and perioperative care in children. Early or primary IO indications are respiratory and circulatory arrest; critical hemodynamic instability before or during anesthesia introduction; severe laryngospasm; anesthesia induction in respiratory bleeding. Urgent indications (decision based on each case is necessary) include urgent induction of anesthesia in non-fasted children (ileus, RSI); induction of anesthesia in children with unstable circulation or severe cardiac insufficiency. Semi-elective indications (decision based on each case is necessary) after mask induction of anesthesia (if vascular access required); mandatory induction of "intravenous" anesthesia (as in malignant hyperthermia). This article is in German.


This article describes an observational study to assess the safety and efficacy of the EZ-IO when using a management algorithm for difficult vascular access in an out-of-hospital setting. Over a one-year period, the device was used in 30 cardiac arrest and 9 other cases. Overall success rate was 97% and first attempt success was 84%. There was one complication—transient local inflammation. Investigators concluded that the device is suitable as a first-line option for difficult vascular access in the out-of-hospital setting.


General overview of PALS updates. Various IO devices were specifically mentioned in the vascular access section, including the EZ-IO.


This article describes a military study in which post-mortem autopsy multidetector CT was used to assess placement of tibial IO needles in battlefield trauma deaths where IO was used as part of the medical intervention. Results showed 58 of 61 (95%) tibial IO needles were correctly placed. In this study, the device used for IO placement was not recorded, but may have been the manual device or EZ-IO as the Army has access to both.


This article summarizes the case-based observations made by the Armed Forces Medical Examiner System on soldiers killed in action/died of wounds who had evidence of sternal intraosseous access. The Pyng Fast-1 is noted in the article as the sternal IO device most widely distributed by the department of defense (DOD); the EZ-IO is listed as another device that may be seen in emergency care facilities within the DOD. Of 98 cases, 78 (80%) showed proper placement; 20% were unsuccessful. It should be noted that the article incorrectly states that the EZ-IO using the powered driver is indicated for sternal placement.

Howarth D. Adult intraosseous access: experiences in a remote emergency department. Australian Family Physician 2011;40(7):510-1

In this article, the author makes a supporting case for remote emergency departments to stock adult intraosseous kits by referencing two adult septic shock cases in which IO access was used for rapid IV fluid replacement as well as IV antibiotics and inotrope support.


This manuscript describes two studies conducted to assess the function and longevity of EZ-IO catheter when placed in the goat model. The authors concluded that the EZ-IO catheter can be left in place for more than 24 hours in animals and can be used in many different veterinary settings when IV access is not immediately available. They also concluded that the EZ-IO system is useful in larger or adult bones.

Khan LAK, Anakwe RE, Murray A, Godwin Y. A severe complication following intraosseous infusion used during resuscitation of a child. Inj Extra 2011;doi:10.1016/j.injury.2011.05.015

This article describes the case of an 11-year-old boy who suffered compartment syndrome of the lower leg following use of the EZ-IO for resuscitation and 24 hours of intraosseous infusion of adrenaline, calcium and potassium. The author concluded that further work is needed to develop recommendations for maximum duration, dose, volume and rates for intraosseous infusion.

This article reports a case in which IO access was used to deliver intravenous contrast agent in an adult blunt trauma patient. After placement in the proximal humerus, contrast agent was administered via the IO route, and clinicians found the CT scans of the thorax, abdomen, and pelvis to be adequate for diagnostic purposes and subjectively equivalent to those of studies using central venous access. There were no complications and the authors concluded that IO catheterization appeared to be an effective alternative to traditional venous access for administering contrast agents for CT evaluation in adult blunt trauma patients.


This abstract describes a swine study presented at the 2011 National Association of EMS Physicians Annual Conference that examined infusion rates through 3 anatomical sites via the powered EZ-IO device. Investigators concluded that the infusion rate was greater via the humerus compared to the tibia and femur.

Larabee TM, Campbell JA, Severyn FA, Little CM. Intrathecal infusion of ice cold saline is less efficacious than intravenous infusion for a CAT scan in a swine model of cardiac arrest. Resuscitation 2011;82(5):603-6. doi:10.1016/j.resuscitation.2011.01.007

This study compared the effectiveness of infusing ice cold saline via IO and IV to induce mild therapeutic hypothermia (temperature drop to 34°C) within a 30 minute timeframe, in a swine model of cardiac arrest. Five swine were evaluated in each the IV and IO groups. Goal temperature was reached in 4/6 animals in the IV group and 0/5 animals in the IO group in the allotted time frame; IV was superior in terms of rate of infusion, rate of temperature change, and time to achieve target temperature.


This study evaluated the use of telesimulation by Canadian pediatricians to teach a relatively new IO insertion technique (EZ-IO System) to physicians in Africa. Self-assessment questionnaires were completed before and after training, multiple-choice tests were given and a demonstration of competency was done within 3 training sessions. Twenty-two physicians participated; the sessions improved physicians’ knowledge, self-reported confidence, and comfort level in inserting the IO needle. The author concluded that telesimulation offers potential for teaching other procedural skills over distances.


This abstract describes an evaluation performed in the goat model, using the EZ-IO, to determine the ability of IO access to accommodate a typical contrast dye infusion and withstand the power injection pressure. Bench testing was done to determine the max pressure deliverable through the EZ-IO using the power injector; various injection occlusion scenarios were established. Results showed the mean pressure through the humerus was 56.5psi; through the tibia was 163.5psi. There were 2 tibial extraosseous distal venous ruptures visible by fluoroscopy but not on gross examination. Under bench testing, for all tests, at pressures up to 750psi no failure or leakage was observed in the IO catheter. The EZ-IO extension tubing should not be used for power injection, particularly if the IO is in the tibia. This research was sponsored by Vidacare Corporation.

Miller LJ, Philbeck TE, Puga TA, Montez DF, Escobar GP. A pre-clinical study to determine the time to bone sealing and healing following intraosseous vascular access. Ann Emerg Med 2011;58(4S):S240

The objectives of this study were to evaluate the amount of time necessary following IO insertion and infusion for the bone to heal such that a second IO catheter can be placed in the same bone without the risk of extravasation from the first hole; and to determine the length of time required to show radiological evidence of closure. Four anesthetized goats were used for the study. Twenty-four hours post insertion, extravasation was observed in 2 of 4 tibial sites with no extravasation in 4 humeral sites. Forty-eight hours post insertion, no extravasation was observed in tibial or humeral sites. Authors concluded that IO infusion should not be attempted in the same bone as a previous IO insertion within 48 hours of removal of the first IO catheter. Radiological examination showed evidence of bone healing as early as 6 days post IO placement. This research was sponsored by Vidacare Corporation.


This article describes the changes in practice experienced when a 12-site statewide ambulance service changed from the manual to the semi-automatic IO device (EZ-IO). There was no statistically significant change in first-attempt success or the number of successes per attempt. However, the use of IO access more than tripled when changing from the manual to the semi-automatic device and PIV access attempts before IO access went from occurring in 35.5% of patients to 1.7% of patients.
Bibliography

Myers LA, Russi CS, Arteaga GM. The introduction of a semiautomated (EZ-IO) device in pediatric prehospital care replacing a manual intraosseous (IO) device improves the success rate for attempts at vascular access. Prehosp Emerg Care 2011;15(1):110

This abstract describes a 93 patient study presented at the 2011 National Association of EMS Physicians Annual Conference that examined the characteristics of pediatric patients receiving IO infusions and the primary EMS clinical impressions, success rates, and subsequent treatments delivered via manual IO vs. the powered EZ-IO device. Investigators concluded that for the pediatric cohort use of the powered device offered a marginally higher first-attempt success rate compared to the manual device; and that the rate of IO access utilization by EMS more than tripled after adoption of the powered device.


This article provides an overview of intraosseous vascular access for pediatrics and discusses general indications, contraindications, complications, and intraosseous devices.


This article in German discusses use of IO access and its multiple applications, focusing on the EZ-IO Infusion System.


This article describes a literature review study with the objective of establishing which intraosseous device is best for prehospital use. This short review searched Medline 1950-2010, CINAHL 1982-2010 and EMBASE 1980-2010 and identified two studies meeting their evidence search criteria, one study compared the BIG vs. manual; the second compared EZ-IO vs. FAST-1. The clinical bottom line asserted by the author was traditional manual IO devices have faster, better success rates in the pre-hospital setting; but that more randomized trials are needed to determine the best device.


This article describes a pre-hospital clinical study comparing IO first-attempt success between humeral and tibial sites. Of 88 cardiac arrest patients analyzed, 58 and 30 IO access attempts were made in the tibia and humerus, respectively. Of those, there was a 90% first attempt success rate for the tibia, compared to 60% for the humerus. Of successful insertions, 6% of tibial insertions became displaced during transport, compared to 33% of humeral insertions. Investigators concluded (the obvious) that "proximal tibial IO needle placement was associated with a significantly higher frequency of first-attempt success and lower incidence of needle dislodgements..." compared to humeral placements.


The objective of this study was to determine the frequency of first-attempt success of humeral IO, tibial IO, and peripheral IV (PIV) insertions during out-of-hospital cardiac arrest. Patients were randomized to receive one of the 3 methods. There were 182 patients enrolled, 64 were assigned to tibial IO, 51 to humeral IO and 67 to PIV. Of all patients 130 (71%) were successful on first attempt with 17 (9%) needles dislodged. First attempt success within the treatment groups was 91% for tibial IO, 51% for humeral IO, and 43% for PIV.


This article describes a case in which systemic fibrinolysis was administered through the intraosseous route in a patient with ST-segment elevation myocardial infarction. Fibrinolytics and antiarrhythmic drugs were administered though the IO line, resulting in resolution of coronary ischemia and electrical instability, without complications. Authors concluded that intraosseous cannulation represents a novel route for administration of systemic fibrinolysis in cases of difficult peripheral venous access in the out-of-hospital setting.


This study conducted in Germany and Switzerland evaluated use of the EZ-IO in the prehospital setting over a 24 month period. The decision to use IO access was left to the discretion of the onsite clinician, a paramedic or an emergency physician. Results showed IO access was attempted in 77 patients, and was successful on first attempt in 75 patients. Significant pain with infusion was reported in the majority of responsive patients.
This document addresses pediatric vascular access and includes an overview of intraosseous vascular access. Indications, contraindications, supplies and equipment, technique, complications and maintenance are discussed.

Sheehan C, Sodhi V, Esler M. Intraosseous needles on the delivery suite. Intraosseous access for neonatal and newborn resuscitation in the national park service (NPS).
This article discusses how a group of obstetricians and anesthesiologists prepared for what they expected to be a difficult delivery with limited venous access. The EZ-IO was brought into the delivery suite as a back-up option if they were unable to achieve venous access in an emergency situation. The authors did note their concern with the pain associated with IO infusion. Ultimately, the IO device was not needed for the delivery in question, but it has been added to their resuscitation kit within the delivery suites.

This article describes IO access in terms of efficacy, indications/contraindications for use, and the IO procedure and comparison of devices to make a case for IO use in oral and maxillofacial surgical practice. In discussing IO devices citing published data, the author identified the EZ-IO device as the most accurate, efficacious, and precise system when trying to achieve IO access.

This article in German concludes that the introduction of IO in pre-hospital pediatric emergency system has markedly reduced the number of critically ill or severely injured pediatric patients without vascular access or with less reliable alternative administration routes in the last 20 years.

This article describes two cases of leg amputation after intraosseous infusion in a 5-month-old girl and a 17-month-old boy. The author concluded that fluid extravasation, exacerbated by tibial fracture and needle dislodgement during transportation, caused limb ischemia in these two patients, and that adherence to the principles of careful needle placement, splinting/securing the catheter and limb, limited length of infusion and repeated monitoring of the limb will help avoid this devastating complication.

This article in Danish discusses use of the IO route for second line vascular access when peripheral IV access is difficult or impossible.

This article is a response to the Taylor and Clarke 2011 report of two amputations required following development of compartment syndrome after IO infusion. The author notes that complications are possible with all methods of establishing IO access including manual, spring loaded and power driven needles and that it is not accurate to directly relate the adverse events to the power driven device only.

YEAR: 2010

This abstract, which was presented at the 2010 ACEP Research Forum, describes a study conducted by investigators from the Medical College of Georgia to determine the frequency of intraosseous vascular access use in adult emergency patients. They surveyed academic emergency departments across the country and, at their own facility, compared ease and speed of standard emergency vascular access methods—including intraosseous. They concluded that IO access is underutilized and generally not the second-line choice of vascular access in unstable adult patients in academic institutions. Their simulation showed IO placement was considerably faster than both central lines and ultrasound guided peripheral IV. They opined that IO should be considered more frequently in critical unstable adult emergency department patients.

This veterinary study evaluated 3 IO access devices, impact driven, automatic rotary, and manual, to compare the placement feasibility and amount of bone trauma induced when used in adult feline cadavers. Seventy-two IO insertion locations were used, the 3 devices were equally randomized to the insertion site. The rotary device was found to have shorter time to insertion and better ease of insertion. No statistically significant differences between number of bone fragments, defect diameter, or success rate were found between devices.
This article reviews intraosseous vascular access and its increased use in adult resuscitation. The IO route is described, including indications, contraindications, insertion sites and devices.

Gillum L. All access pass: mastering the use of IO devices. JEMS 2010;35(6):30,32
This article discusses training methodology and applies the concept to the implementation of the EZ-IO in the Montgomery County Hospital District EMS, a participant in the EZ-IO beta test.

An article evaluating various methods of obtaining vascular access in the management of 21st century battlefield trauma including, peripheral IV access, intraosseous access, venous cut-down, and central venous access. The authors conclude that IO access should be the first line vascular access in casualties with severe trauma to avoid delay initiating resuscitation in pre-hospital or hospital setting.

This article in German describes the results of a survey of rescue assistants and physicians, in which they found that IO use was still a rarity in the Berlin emergency medical service and, therefore, presumably nationwide.

Authors describe an early observational study (N=120) comparing intraosseous access in the humerus and the tibia, using the EZ-IO. Investigators concluded that the humerus is an acceptable IO site, which may be preferable under certain clinical conditions. This research was sponsored by Vidacare Corporation.

In an abstract presented at the 2010 ACEP Research Forum, investigators describe a swine study designed to compare IO infusion rates using the Belmont FMS 2000 rapid infusion device and a pressure bag through the proximal tibia and proximal humerus. Investigators concluded that infusion rates were highest using the pressure bag via the proximal humerus.

Authors describe a randomized, controlled trial comparing two different IO access devices in adults in the hospital setting. Twenty patients received the BIG and 20 received the EZ-IO. Success rate on first attempt was 80% for the BIG and 90% for the EZ-IO. Mean procedure time was 2.2 minutes for the BIG vs. 1.8 minutes for the EZ-IO. Differences in success rate and procedure time were not statistically significant, and there were no significant complications for any patients. Investigators concluded that IO access is a reliable and safe method for rapid vascular access for in-hospital adult patients under resuscitation.

This article provides an overview of intraosseous vascular access and discusses general indications, contraindications, complications, and intraosseous devices.

In this study, using a swine model, investigators concluded that mild therapeutic hypothermia can be effectively induced after successful resuscitation of prolonged ventricular fibrillation through infusion of chilled saline via the IO catheter.


This abstract, presented at the 2010 ACEP Research Forum, describes study designed to determine the relative precision of intraosseous needle placement using only tactile feedback. The study also assessed the ability to access simulated osteoporotic bone without damage using the 3 methods. Investigators concluded that, using tactile feedback only, rotary power may allow precise IO catheter placement with greater success and confidence than manual or hammer-assisted devices. Powered insertion may facilitate penetration into fragile bone without damage. This research was sponsored by Vidacare Corporation.


This abstract, presented at the 2010 ACEP Research Forum, describes a study designed to determine infusion flow rates through the proximal humerus and proximal tibia. Investigators found that, at all infusion pressure levels, the humerus provided substantially greater flow rates than the tibia. They concluded that, for most situations, adequate IO infusion rates can be achieved using the tibial site, but the proximal humerus site should be strongly considered when greater infusion flow rates are required. This research was sponsored by Vidacare Corporation.


This abstract presented at the 2010 ACEP Research Forum describes a study designed to compare Lidocaine’s effect on pain during fluid infusion through the tibial and humeral IO routes. Authors concluded that, for adequate IO infusion rates with minimal and tolerable pain, 40mg of preservative-free Lidocaine may be needed; followed by a rapid normal saline syringe flush of at least 10mL and another 20mg of Lidocaine. Additional dosing and flushing may be required. For less overall pain due to IO infusion, and greater infusion flow rates, the proximal humerus should be strongly considered, using a longer IO needleset. This research was sponsored by Vidacare Corporation.


Authors describe a 10 subject volunteer study that compared intraosseous (IO) blood samples to venous blood samples for complete blood count (CBC) and chemistry profile testing. They concluded that IO blood may serve as a reliable alternate for hemoglobin and hematocrit levels, as well as for most analytes in a basic blood chemistry profile. Exceptions are CO₂ levels, platelets, and WBC. This research was sponsored by Vidacare Corporation.


In this abstract of a study presented at the 2010 National Association of EMS Physicians Meeting, researchers describe a study in which sternal and tibial IO devices were evaluated with and without chemical protective equipment. Researchers concluded that the use of the protective equipment did not affect the success rate or time to placement for the two IO devices.


This article describes an online questionnaire study in which the Heads of Department of 20 EDs in Denmark were asked about IO infusion within their institution. Nineteen responses were received; 14 hospitals (74%) reported having IO devices available with the median number of IO procedures performed as 5. In 9 departments training had not been provided and 8 departments didn’t have IO guidelines. The favored device was the EZ-IO found in 18 of the EDs, 2 had EZ-IO and Cook Surfast and 1 had the BIG.


Authors report an observational study of 14 children in whom semi-elective IO infusion was performed under anesthesia after peripheral IV had failed. IO infusion was successful for all 14 patients, using the EZ-IO system for 8 patients and the Cook system for 6 patients.
Philbeck TE, Miller LJ, Montez D, Puga T. Hurts so good; easing IO pain and pressure. JEMS 2010;35(9):58-69

This article describes two studies designed to compare Lidocaine’s effect on pain during fluid infusion through the tibial and humeral IO routes and to determine infusion flow rates. Authors concluded that, for adequate IO infusion rates with minimal and tolerable pain, 40mg of preservative-free Lidocaine may be needed; followed by a rapid normal saline syringe flush of at least 10mL and another 20mg of Lidocaine. Additional dosing and flushing may be required. For less overall pain due to IO infusion, and greater infusion flow rates, the proximal humerus should be strongly considered, using a longer IO needleset. This research was sponsored by Vidacare Corporation.


A simulation study evaluating if use of a laryngeal mask airways (LMA) and intraosseous (IO) lines established using the EZ-IO leads to improved resuscitation in a simulated cardiac arrest when compared to standard endotracheal intubation and central line placement. Results showed mean time to airway, mean duration of airway attempt, and time to vascular access was shorter in the IO group than the CVL group. Time to defibrillation and percent hand off time was not significantly different between the groups.


This abstract describes an animal study, presented at the 2010 ACEP Research Forum, that examined shear and pressure changes within the medullary space during intraosseous infusion. Results suggest that resistance to flow depends on cannula placement site, IO pressure rises rapidly with infusion rates, and medullary compression and axial shear are present at high infusion rates.


This letter to the editor is written in response to the case report by Landy titled, &quot;Complication of intraosseous administration of systemic thrombolysis for a massive pulmonary embolism with cardiac arrest.&quot; The author suggests that the tissue necrosis described by Landy may have been due to the removal of the IO needle while there was still significant fibrinolytic activity at the needle insertion site. The author suggests a change in medical care after return of spontaneous circulation (ROSC) in patients following thrombolytic administration through IO access to convert the functioning IO line to a non-flowing saline lock. The EZ-IO was used to provide IO access in the case report by Landy.


This article describes a longitudinal study of intraosseous vascular access in pre-hospital emergency medicine handled by helicopter emergency medical services. Of the 78 IO insertion attempts made on 70 patients, overall success rates were 50% using manual needles, 55% using the Bone Injection Gun, and 96% using the EZ-IO. Investigators concluded that newer IO techniques may enable faster and more reliable vascular access; and that all emergency services should be familiar with IO techniques.

Tobias JD, Ross AK. Intraosseous infusions: A review for the anesthesiologist with a focus on pediatric use. Anesthesia & Analgesia 2010;110(2):391-401

The authors describe literature that support the use of IO access for administering anesthesia in the ICU, perioperative and operating room, including a study in which IO access was used successfully for providing intraoperative anesthesia for 106 of 109 patients. Among their conclusions, the authors reported that, even though rarely reported in anesthesia literature, IO access is a technique anyone providing care to children should consider when the patient has difficult IV access. They also concluded that IO access should be a part of an algorithm that includes numbers of attempts at peripheral access, feasibility of central access and the need for continued postoperative access; and that considering that IO access may be occasionally used in the perioperative setting in both emergent and nonemergent scenarios, it may be beneficial to have appropriate IO needle in the OR.

Turkan H. How does the training effect the use of intraosseous access with a battery driven device?. Resuscitation 2010;81(2):S93. doi:http://dx.doi.org.10.1016/j.resuscitation.2010.09.380

This abstract describes a study in which 60 physicians, nurses, and paramedics naïve to IO vascular access were trained on the Arrow EZ-IO system. After lecture and hands-on training, the clinicians attempted to perform the procedure using a bone model and evaluated the device for ease of insertion, number of attempts, time to insertion, and their opinion on the device. The authors concluded use of the EZ-IO system can result in high success rates of insertion with inexperienced device users.

Vegunta RK. Chapter 8-Vascular access. Ashcraft’s Pediatric Surgery 2010;5th ed:110-16

This document discusses various vascular access methods available for pediatric and neonate patients, including intraosseous access.
Vizcarra C, Clum S. Intraosseous route as alternative access for infusion therapy. Journal of Infusion Nursing 2010;33(3):162-74
This article provides an overview of IO anatomy and physiology, IO access indications, care, and management; describes therapies administered via IO access; and discusses the expanding use of IO access into areas within hospitals during nonemergency clinical situations. It also includes a table addressing indications for IO access in the hospital, as well as a table addressing the general insertion procedure for IO access.

This retrospective study evaluated humeral IO placement success rates, using the EZ-IO, in the out of hospital cardiac arrest patient. Over a 9 month period, humeral placement was attempted in 247 patients. First attempt successful placement was 91%; successful placement within two attempts was 94%. The authors concluded that humeral IO is a reliable method of fluid and drug delivery in the out of hospital cardiac arrest population. This research was sponsored by Vidacare Corporation.

YEAR: 2009

This study was designed to evaluate the effect of education on knowledge, attitudes and skill performance of IO access by Level 1 EMTs in Korea. After a two-hour program, the knowledge and attitude of IO access improved significantly.

In a goat study, researchers assessed the hemodynamics of hydroxocobalamin (OHCo) and normal saline (NS) by the IO route and concluded that the effects of OHCo given by the IO route in non-CN-poisoned goats are mild and well tolerated.

This letter to the editor discussed the experience of one ground emergency rescue service in Germany and their trial implementation of the EZ-IO, as compared to the David et al evaluation of the BIG by emergency physicians in which the rate of failure was 55%. Over a one year evaluation of the EZ-IO in the field, it was used in 20 patients, of which 19 were successfully placed (95%). The success of the field evaluation and a human cadaver study resulted in the addition of the EZ-IO to the receiving University Hospital emergency department.

Case report of IO infusion in 79-year old woman with hematemesis after intestinal surgery.

This article provides a general overview of intraosseous access and its use in emergency situations. A description of available IO access devices is provided.

Prospective study of 246 EMS providers at 14 EMS agencies. Reports successful IO placement in 95% of cases (18 of 19).

This article describes a study conducted at an urban Level I trauma center in Munich, Germany. Ten consecutive patients for whom PIV was difficult or impossible were simultaneously given a central line and an EZ-IO. Procedure times were measured and defined as the time the device package was taken off the shelf until the first drug or solution was administered. First attempt success rate was 90% for EZ-IO and 60% for CVC. The mean procedure times were 2.3 minutes for EZ-IO and 9.9 minutes for CVC, a clinically and statistically significant difference. Investigators concluded, because CVC was "slower and less efficacious..." IO may improve the safety of patients requiring resuscitation in the ED.
This article describes a cadaver study to determine skill acquisition and performance by use of the EZ-IO system by novices. Overall success rate for the 99 operators was 97%, and mean insertion time was 6 seconds. All operators rated the device faster and easier than using a central line, and 99% expressed willingness to use the device for cardiac arrest patients.

This article describes IO use in general, and the EZ-IO in particular. The author describes its use by the emergency staff at her hospital and how they became advocates for IO access in both emergent adult and pediatric patients. She found that its use improves the quality of our care by providing vascular access to our most critical patients.

This abstract for a presentation at the 2009 ACEP Research Forum describes a swine study designed to determine the feasibility of inducing therapeutic hypothermia (TH) after resuscitation by giving an IO infusion of iced saline. Researchers concluded that rapid, large volume IO infusion of iced saline is as effective for lowering core body temperature after resuscitation as central access and peripheral IV. This research was sponsored by Vidacare Corporation.

This abstract describes a retrospective study to determine the time from EMS dispatch to IV or IO drug delivery, time savings to drug delivery if vascular access preceded intubation, the internal validity of that point estimate using matched cases in which IV/IO was performed first, and the theoretical increase in rate of return to spontaneous circulation. Investigators concluded that time from dispatch to IV/IO delivery could be reduced by 4 minutes if vascular access preceded intubation and could, potentially double ROSC.

This abstract for a presentation at the 2009 ACEP Research Forum describes a swine study that evaluated crystalloid fluid flow through an IO needle following nitroglycerin infusion in a swine model. Investigators concluded there was not a significant increase in flow rate after administration of IO nitroglycerin.

This abstract for a presentation at the 2009 ACEP Research Forum describes a volunteer study that examined the relationships between IO and venous blood samples when analyzed for complete blood count and chemistry profile. Researchers concluded that the IO space is a reliable source for blood used for CBC and chemistry profile. Results may be moderately reliable for carbon dioxide, but unreliable for WBC counts that appear to be elevated and platelet counts that appear lower. This study was sponsored by Vidacare Corporation.

Miller LJ, Philbeck TE, Montez DF, Spadaccini CJ. A new study of intraosseous blood for laboratory analysis. Arch Pathol Lab Med 2009;133:1628
This abstract for a presentation at the College of American Pathology 2009 meeting describes a volunteer study that examined the relationships between IO and venous blood samples when analyzed for complete blood count and chemistry profile. Researchers concluded that the IO space is a reliable source for blood used for CBC and chemistry profile. Results may be moderately reliable for carbon dioxide, but unreliable for WBC counts that appear to be elevated and platelet counts that appear lower.

This article describes a prospective, observational study involving a convenience sample of 25 medical students, physicians and nursing staff recruited evaluate the EZIO powered drill device on a bone model. Twenty-three (92%) of the 25 study subjects required only one attempt at placing the EZ-IO. Investigators concluded that the device was easy to use with high success rates of insertion with inexperienced participants.
This article describes a case in which IO access, using the EZ-IO, was attempted in a patient with osteogenesis imperfecta. In each of 3 attempts, the needle became loose immediately after IO insertion. The author acknowledged that during emergencies it is difficult to assess and consider every possible contraindication for every intervention; and that IO access using the EZ-IO is the author’s choice for emergency vascular access when peripheral access is difficult or has failed.

Ong ME, Chan YH, Oh HH, Ngo AS. An observational prospective study comparing tibial and humeral intraosseous access using the EZ-IO. Am J Emerg Med 2009; 27: 8-15
Comparison of tibial and humeral IO use in 24 adults. Both sites suitable for IO infusion. This study was sponsored by Vidacare Corporation.

This article describes a prospective, observational study involving a convenience sample of 25 medical students, physicians and nursing staff recruited to evaluate the EZIO powered drill device on a bone model. Twenty-three (92%) of the 25 study subjects required only one attempt at placing the EZ-IO. Investigators concluded that the device was easy to use with high success rates of insertion with inexperienced participants. (Note: This study was also described in an earlier article published in American Journal of Emergency Medicine) This study was sponsored by Vidacare Corporation.

In this 1,598 patient case series, investigators studied the effects of "serial standard of care changes...in the EMS system over time." They concluded that IO access is an essential component for a proven algorithm for the management of OOH-CA.

This article describes the first clinical study that focuses on the proximal humerus as an IO site. It is also the first article describing a comparison between IO access and peripheral IV (PIV) and central venous catheters (CVC). They found that IO catheter placement was significantly faster than PIV or CVC placement, and concluded that IO access is life-saving when PIV or CVC is difficult or impossible. This study was sponsored by Vidacare Corporation.

This abstract for a presentation at the 2009 ACEP Research Forum describes a volunteer study to determine the optimal Lidocaine dosing and sequencing for patients receiving fluids through the IO route and to determine if adequate fluid flow rates can be delivered through the proximal humerus IO site. Researchers concluded that for adequate IO infusion rates with minimal and tolerable pain, 40mg of preservative-free Lidocaine may be needed, followed by a rapid normal saline flush of 10ml. Additional dosing and flushing may be required. For humeral insertion, a longer IO needleset should be considered. This study was sponsored by Vidacare Corporation.

This article describes an observational study performed by the French military air surgical team in Chad. There were 11 patients with no insertion failures. For 7 patients, the insertion site was the proximal tibia and for the remainder the site was the proximal humerus. The authors concluded that the EZ-IO is a device that is simple, reliable and which gives satisfaction for the administration of drugs.

This case study describes injuries sustained in Iraq by an American soldier, and the concurrent use of 4 IO devices to resuscitate and stabilize him.

This abstract describes a small study designed to determine if IO line placement improves outcome in adult patients with out-of-hospital cardiac arrest. This 165 patient study did not demonstrate improved survival.

Authors of this article describe a pilot study designed to compare the success rate for insertion and ease-of-use of the Bone Injection Gun (BIG) spring-loaded device and the EZ-IO battery-powered device on a turkey bone model. Investigators concluded that the EZ-IO demonstrated higher success rates than the BIG (28/29 vs. 19/29, p=0.016), and the EZ-IO was the preferred device.


This article, in German, describes the technique of IO access, the introduction of two different IO devices (Cook and EZ-IO) and describes IO use in pediatric emergency care.


This review article in German describes intraosseous vascular access, and includes descriptions of the Waismed Bone Injection Gun, Vidacare EZ-IO, Jamshidi and Cook Medical IO devices.


This letter to the editor describes the first case reported in the clinical literature in which therapeutic hypothermia was administered using the intraosseous route. The patient, a 2-year-old boy who was found submerged in a cesspool and had been asystolic for 5-10 minutes, survived without neurological complications

Von Hoff DD, Kuhn JG, Burris HA, Miller LJ. La perfusion intraosseuse est-elle equivalente a la perfusion intraveineuse?. Urgence Pratique 2009;36:36-40

This French version of an article previously published in American Journal of Emergency Medicine describes a 25-patient clinical study that compared the pharmacokinetics of intraosseous using the VidaPort (a predecessor of the Vidacare EZ-IO) vs. intravenous administration of morphine sulfate in adults. Results showed no differences between IO and IV administration of morphine for nearly all pharmacokinetic parameters, including maximum plasma concentration, time to maximum plasma concentration, and area under plasma concentration-time curve. There was a significant difference in the volume of distribution in the central compartment, which investigators attributed to a minor deposition effect near the IO port or in the bone marrow. Investigators concluded that the results support the bioequivalence of IO and IV administration of morphine in adults.


This article, in Norwegian, describes IO access and modern IO devices, including the Bone Injection Gun, FAST1, and EZ-IO.


This swine study was designed to determine if intraosseous infusion is suitable to delivery recombinant human factor VIIa (rFVIIa) during hemorrhagic shock. Investigators concluded that administration of rFVIIa via IO infusion is a safe route for delivery and is likely to produce blood levels required to improve hemostasis during shock.


This article discusses the importance of proper technique, attention to detail, and serial monitoring of limb involved when using IO vascular access to avoid potential compartment syndrome and other complications. The author reports the case of a 2-year-old boy who suffered compartment syndrome of the lower limb following use of IO infusion for resuscitation. Early detection of and response to changes in the affected limb resulted in the patient’s successful recovery


Animal (goat) study to determine if IO administration of hydroxocobalamin for antidotal treatment for exposure to cyanide and other poison agents would be faster and require less fine motor coordination and sensitivity; and would result in similar hemodynamic changes compared to IV administration. Using the EZ-IO device, researchers concluded that hemodynamic effects of hydroxocobalamin given by the IO route in non-poisoned goats are mild and similar in magnitude to those of saline control animals.

Animal (goat) study to determine the capacity and time required for protected hazardous materials responders and receivers to accomplish vascular access and hydroxocobalamin administration for antidotal treatment for exposure to cyanide and other poison agents. Using the EZ-IO device, researchers concluded that the time required for IO administration of the drug was shorter than intravenous administration; and that IO placement is readily accomplished wearing all levels of chemical protective garments and equipment.


Study comparing manual intraosseous insertion with EZ-IO using adult human cadavers as a model. No significant difference in insertion time between 39 manual insertions and 45 EZ-IO insertions. Found a difference in the success rate (manual, 79.5% vs. EZ-IO 97.8%, p<0.01). The EZ-IO had fewer complications (manual, 15.4% vs. EZ-IO 0.0%, p<0.01) and scored higher on user friendliness (school grading system: manual, 1.9±0.7 vs. EZ-IO 1.2±0.4, p<0.01).


Describes common drugs used in pediatric resuscitation and evidence supporting their use. Also describes routes of administration including intravenous, intraosseous, and intratracheal. Describes IO systems including Bone Injection Gun, FAST-1, and EZ-IO.


This article describes IO infusion devices - including Jamshidi, Cook, WaisMed, and Vidacare devices - and placement sites. It also addresses assessment and care of the infant receiving fluids and medications through the IO route.


Large retrospective study of patients for whom emergency vascular access was obtained using the Vidacare EZ-IO intraosseous system. Insertion success was 92% and within 10 seconds for 84% of the one-attempt successful cases. Complication rate was low (4.8%), none were serious, and extravasation was the most frequent (0.8%). The device was rated easy to use 72% of the time, and researchers concluded that the powered IO device is safe and effective for achieving vascular access in the resuscitation and stabilization of emergency patients. This study was sponsored by Vidacare Corporation.

Horton MA, Beamer C. Powered intraosseous insertion provides safe and effective vascular access for pediatric emergency patients. Pediatr Emerg Care 2008;24:347-50

A retrospective clinical study was conducted to demonstrate the safety and effectiveness of the EZ-IO intraosseous access device for pediatric patients. For the 95 eligible patients in the study, successful insertion and infusion was achieved in 94% of the patients. Insertion time was 10 seconds or less in 77% of the one-attempt successful cases reporting time to insertion. There were 4 minor complications (4%), but none significant. The results of this study support the use of the EZ-IO for children in emergency situations. The complication rate suggests that the powered IO device is safe and effective for the resuscitation and stabilization of pediatric patients. This study was sponsored by Vidacare Corporation.


Article describing IO access.


Interim report for quasi-controlled prospective study of emergency department patients for whom emergency vascular access using the Vidacare EZ-IO intraosseous (IO) system (n=6) inserted in the proximal humerus was compared to access using central or peripheral intravenous (IV) lines (n=60). Researchers concluded that proximal humerus IO insertion is significantly faster than central or peripheral intravenous (IV) line insertion. Complications and pain profiles were similar for IO and IV techniques. This study was sponsored by Vidacare Corporation.
This article describes an observational study in which two intraosseous devices were compared: the Pyng Medical F.A.S.T.1 and the Vidacare EZ-IO. For the 117 patients on which the F.A.S.T.1 was used, there was an 84% success; compared to a 97% success rate for the EZ-IO (n=71).

This article describes a 25-patient clinical study that compared the pharmacokinetics of intraosseous vs. intravenous administration of morphine sulfate in adults. Results showed no differences between IO and IV administration of morphine sulfate for nearly all pharmacokinetic parameters. Investigators concluded that the results support the bioequivalence of IO and IV administration of morphine in adults.

This article describes thoracic trauma in the pediatric population. Includes a review of the assessment of pediatric patients. Circulation section of the article strongly recommends rapid intravascular volume expansion by the intraosseous route, and recommends the EZ-IO for "...quick and reliable vascular access during resuscitation ...".

**YEAR: 2007**

This abstract for a presentation at the 2007 American College of Emergency Physicians Research Forum describes an observational study in which the EZ-IO was used to provide emergency vascular access for 95 pediatric patients. Successful insertion and infusion was achieved in 94% of the patients, and insertion time was within 10 seconds for 81% of the placements. There were four minor and no serious complications.

This article reviews and assesses the literature on the use of IO drug administration during cardiopulmonary resuscitation. It addresses the risks and benefits of using IO in adults and children. The article describes the FDA-cleared devices available for use including the Pyng F.A.S.T.1, Waismann Bone Injection Gun and the Vidacare EZ-IO.

Describes the experience of the UK Defence Medical Service using the EZ-IO for emergency vascular access in Afghanistan. They used the device for 26 patients, including 10 children. Of the 26 EZ-IO placements, 23 were made in the emergency department. There was a 97% insertion success rate with no infection. Significant infusion pain was felt by three patients.

de Caen A. Venous access in the critically ill child. Pediatr Emerg Care 2007;23:422-4
This review article states the availability of intraosseous (IO) needles for pediatric patients, outlines the limitations of traditional venous access, and discusses the various IO devices currently available, including the Vidacare EZ-IO®.

This article summarized the challenges and methods of providing vascular access for infants. It describes IO techniques and devices, including the Jamshidi, Cook, EZ-IO® and Bone Injection Gun (BIG) devices.

Article calls for EMS medical directors to consider and use the intraosseous route for adult patients requiring immediate vascular access. Provides evidence in support of position statement by the National Association of EMS Physicians on IO use.
This article describes authors' evaluation of provider performance using two IO devices; the Pyng Medical F.A.S.T.1™ and the Vidacare EZ-IO®. Of 89 insertions with each device, success rate for 72% for the F.A.S.T.1 and 87% for the EZ-IO, a significant difference ($p=0.009$). The time to fluid insertion for the EZ-IO was also faster ($p=0.02$). Authors noted that the EZ-IO is unique and much more useful than the F.A.S.T.1.

Article in Italian describing IO access and EZ-IO

This abstract for a presentation at the 2007 American College of Emergency Physicians Research Forum describes an observational study done at Boston Medical Center in which the Vidacare EZ-IO was used to provide emergency vascular access for 50 critically-ill adult patients. Successful insertion was achieved in 92% of the patients; with 90% success on the first attempt. There was one immediate complication—a dislodgement during transport. Investigators concluded that the device is a safe and feasible device for adult patients requiring out-of-hospital vascular care.

Results from this, study which sought to compare drug delivery time using the proximal humerus IO route to delivery time using the sternal IO route, suggest that IO proximal humerus is comparable to IO sternal for prompt drug delivery during CPR.

Overview of IO access. Includes historical aspects, current status, indications for use, advantages and disadvantages, IO kinetics, insertion sites, complications and contraindications and description of available IO devices, including EZ-IO®.

In this study, presented at the NAEMSP 2007 annual meeting, authors compared the success rate of conventional IO access with the EZ-IO during 245 cases in the prehospital setting. They concluded that using EZ-IO® results in a statistically significant increase in IO success rate, compared to conventional IO methods.

Myers BJ, Lewis R. Induced cooling by EMS (ICE): year one in Raleigh/Wake County. JEMS 2007; 32: s13-5.
This article describes the experience of the Wake County (NC) EMS System in inducing hypothermia for patients with return of spontaneous circulation after cardiac arrest. Authors describe their use of the Vidacare EZ-IO for the administration of chilled saline; with 414 placements and an overall success rate of 94%.

Potyka JS, Gordon DJ. Stories behind the numbers: IO experiences in providers’ own words. JEMS 2007; 32: s30-1.
Qualitative study focuses on EMS caregivers’ experiences with Vidacare’s EZ-IO device and personal opinions. The study used a narrative approach to gain insight from EMS practitioners working with an IO access device under real field conditions.

Pye D. NY Paramedics get the EZ-IO. Journal of Emergency Medical Services 2007; doi: http://www.jems.com/print/5184
This article in JEMS discusses an EMS system in New York following their adoption of the EZ-IO, and the advantages.


The article describes a prospective observational study conducted by several EMS agencies in Portland, OR to determine the safety, efficacy and benefits of using the Vidacare EZ-IO in the prehospital environment. The IO device was successfully placed in 95% of the 280 cases. In 98% of the cases, placement was made within six seconds.


Study investigating time difference in obtaining IO vs. IV access while wearing personal protective equipment (PPE) in simulated HazMat scenarios. With provider in PPE and mannequin not in PPE, vascular access was faster with IO (14 seconds vs. 46 seconds, p&lt;0.001); also, fluid infusion time (28 seconds vs. 46 seconds, p&lt;0.001). With provider and mannequin in PPE, all the following favored IO: needle to skin time (13 seconds vs. 25 seconds, p&lt;0.001), vascular access time (17 seconds vs. 63 seconds, p&lt;0.001), and fluid infusion time (30 seconds vs. 66 seconds, p&lt;0.001). Investigators conclude that EZ-IO under HazMat conditions provides vascular access and fluid more quickly than IV access.


Article describes a controlled study in which the time difference between IV and IO access was compared while providers and simulated patients (mannequins) were wearing personal protective equipment (PPE). Twenty-two EMT-P providers measured the times to skin access, vascular access and fluid infusion in three scenarios: no PPE for providers or mannequins; providers only in PPE; and both providers and mannequins in PPE. In all scenarios, there was a statistically significant difference in vascular access and fluid infusion time, in favor of the EZ-IO. Investigators concluded that, overall, the EZ-IO provides vascular access and fluid more quickly than standard IV access, and that donning PPE does not hinder providers’ use of the EZ-IO.

Wayne MA. Intraosseous vascular access: devices, sites and rationale for IO use. JEMS 2007; 32: s23-5.

This article reviews intraosseous vascular access in general, and summarizes the various devices available. These include the Waismed B.I.G., the Vidacare EZ-IO, and Pyng F.A.S.T.1.


This article in German (with abstract in English) describes IO infusion in detail. It includes techniques, indications, complications, and recommendations. Also describes the various devices available, including Cook, Bone Injection Gun (BIG), First Access for Shock and Trauma (F.A.S.T.1), and the EZ-IO®.

YEAR: 2006


In this study, presented at the NAEESP 2006 annual meeting, investigators reported the results of a study that evaluated the performance of the EZ-IO® compared to an earlier evaluation of the Pyng F.A.S.T.1 system. There was a statistically significant higher success rate using the EZ-IO® compared to Pyng system, and investigators concluded that the EZ-IO® appears to be a superior device with regard to insertion success.


Prospective observational study evaluating EMT-B ability to provide care in out-of-hospital cardiac arrests. Found that EMT-Bs were able to place the EZ-IO with a 94% success rate. Median time to placement was 72 seconds.

Hoskins SL, Kramer GC, Stephens CT, Zachariah BS. Efficacy of epinephrine delivery via the intraosseous humeral head route during CPR. Circulation 2006;114:I_1204

Results from this study which sought to determine the efficacy of intraosseous drug delivery using the proximal humerus during CPR in swine showed that the humeral route generated higher mean arterial pressures than central venous or endotracheal delivery.

Article in French describes IO access and IO devices, including B.I.G., F.A.S.T.1 and EZ-IO®.


The author provides an overview of intraosseous vascular access discussing evolution of the practice, equipment, treatment options and contraindications.

**YEAR: 2005**


Observational study evaluating use of the EZ-IO®. Found 97% success rate for insertion and infusion into the IO space by paramedics, nurses, physicians and other EMS personnel in using the device for emergency vascular access. No serious complications reported. This study was sponsored by Vidacare Corporation.


Article describes intraosseous access for adults and pediatrics. Describes and discusses IO devices available including Jamshidi, Bone Injection Gun, F.A.S.T.1, and EZ-IO®.


Observational study of initial use of the EZ-IO® in 125 patients by EMS providers. Found 94% success rate for insertion and infusion into the IO space. No complications reported. This study was sponsored by Vidacare Corporation.

Heightman AJ. The rebirth of adult IO: a first-hand account of recent advances in intraosseous infusion for adults, drawn from a scientific workshop and practical lab experience. JEMS 2005;30(10):s4-7.

Editorial article highlighting recent advances in intraosseous (IO) infusion and IO devices based on the author’s experience at a scientific seminar hosted by Vidacare. Makes recommendations on the efficiency and safety of the devices.


This animal study compared IO drug delivery in the tibia versus the sternum during CPR. Researchers concluded that during CPR IO infusions delivered via both sites were effective—although sternal delivery was faster; and that IO sternum access is comparable to IV access for drug delivery during CPR.


This study abstract discusses use of the EZ-IO to determine the pharmacokinetics (PK) and efficacy of tibial IO drug delivery during treatment of cardiac arrest in the swine model, as compared to IV access. Results showed that PK analysis of IO drug delivery via the tibial route showed a delay of 20-50 seconds compared to IV; however, physiologically significant levels of epinephrine were reached as MAP. This research was sponsored by Vidacare Corporation.


Animal study compared the sternal and tibial routes for IO drug delivery during CPR. Investigators concluded that both the sternal and tibial routes can effectively deliver near equivalent doses during CPR in swine. http://www.aemj.org/cgi/content/abstract/12/5_suppl_1/67

Miller LJ, Kramer GC, Bolletter S. Rescue access made easy: Intraosseous infusion, once limited to use in children, is now becoming a reliable access site for adults. JEMS 2005;30:s8-18.

Overview of IO therapy. Includes &quot;10 Myths about Adult IO,&quot; and description of available IO devices, including EZ-IO®.

Seminal article on blood circulation in the IO space. Demonstrates movement of red blood cells from the bone marrow into the circulating blood by perfusion of the tibia of the dog and by injections into the bone marrow in the rabbit and cat.