

# HURON PERTH HEALTHCARE ALLIANCE MEDICAL DIRECTIVE

Medical Directive	Intraosseous Insertion –Paediatric
Directive #	MD-ED-CC-028
Approval	Medical Advisory Committee
Date	November 22, 2018
Signature	Janu moore
Review/Revision Date	Original Oct/18
Specific to	HPHA Critical Care Unit and Emergency Medicine Departments

## **Description of Procedure:**

Insertion of an Intraosseous (IO) needle into the bone marrow using a battery-powered hand held drill in the pediatric patient requiring immediate emergency administration of medications and/or fluids.

If the pediatric patient is responsive to pain and when doing so would not delay emergent treatment, consider administering 3mg/kg of Lidocaine 1% (10 mg/mL) subcutaneously to the identified site and allow 1 minute for effect, prior to IO insertion.

Example for a child weighing 10kg: the dose is 30mg, which is 3mL of a 1% (10mg/mL) solution

If the patient is responsive and patient condition allows, consider administering Lidocaine prior to flushing the IO post insertion:

<u>Pediatrics</u>: inject 0.5 mg/kg preservative free Lidocaine 2%(20 mg/mL) to a maximum of 60 mg (3mL) via IO over 120 seconds, allow Lidocaine to dwell in IO space 60 seconds, then flush with 2-5 mL Sodium Chloride 0.9%.

An intraosseous (IO) needle should be removed as soon as reliable peripheral or central venous access is obtained. An IO needle should not remain in place for more than 24 hours.

#### **Authorized To:**

Registered Nurses employed in the Critical Care Unit and Emergency Medicine
Departments at the Huron Perth Healthcare Alliance who have successfully obtained
certification specific to this particular skill are eligible to implement this directive.

## **Specific Patient Conditions:**

- Any pediatric patient at the Huron Perth Healthcare Alliance who is responsive to pain, in a potentially life threatening situation when IV access has been unsuccessful in 2 attempts or 90 seconds.
- Any pediatric patient experiencing full cardiac arrest or severe shock should have IO cannulation considered as the initial method of obtaining vascular access.

#### **Contraindications:**

- Non emergent situations where IV access can be readily obtained.
- Patient capable of consent refuses treatment or substitute decision maker refuses on behalf of the patient.

## Note: If a patient or substitute decision maker refuses treatment contact the physician immediately to determine plan of care.

- Any documented allergy to Lidocaine is a contraindication to the administration of lidocaine only.
- Fracture of the tibia or femur (consider alternate tibia)
- Previous significant orthopedic procedure
- Previous IO attempt in the same bone within 48 hours
- Pre-existing medical condition affecting the integrity of the bone (tumor near site, osteogenesis imperfecta, etc)
- Infection at insertion site (consider alternate site)
- Inability to locate landmarks (significant edema)
- Excessive tissue at insertion site (obesity)

## Reasons to seek immediate medical consultation or discontinue procedure/ treatment/intervention:

- Patient capable of consent refuses treatment or substitute decision maker refuses on behalf of the patient.
- Through-and-through penetration of the bone
- Occlusion of the IO needle
- Change in sensation, pulses, or perfusion distal to the IO needle
- Signs of infiltration such as increased fluid leakage around the insertion site, new edema
  or increased diameter of the accessed extremity

#### **Documentation:**

 Implementation of the Medical Directive including name and number of the directive, name, signature and credentials of the implementer and name of the attending physician in the order section of the chart.

## **Quality Assurance**

- The Medical Program Director, Emergency Medicine and/or Medical Program Director, Medicine will approve the education component of the Medical Directive.
- The Emergency Department/Critical Care Registered Nurse will complete an annual educational component specific to this medical directive as outlined in the related HPHA Policy to be eligible to implement the directive.
- The Emergency Department/Critical Care Registered Nurse will demonstrate competence in the Medical Directive prior to initiating.
- An annual review will be conducted at the discretion of the HPHA Emergency Care Team to review the appropriateness of the Medical Directive.

Originator	HPHA Clinical Nurse Educators
Current Review/Revision	October 2018
Responsibility HPHA Joint Emergency/Critical Care Team	
Distribution	HPHA Emergency Department Manuals/Critical Care Unit Manuals HPHA My Alliance CCU/Telemetry and ED Medical Directives HPHA Start Hub link ED Medical Directives

## **Appendix**

For step-by-step review of anatomical site identification Click here

## Reference(s):

Elsevier Performance Manager (2018). Clinical Skills: Intraosseous Needle Placement (Pediatric)

Lakeridge Health. (2012). Medical Directive-Advanced Life Support: Intraosseous (IO) Cannulation and Infusion.

Saskatoon Health Region. (2011). Intraosseous Infusion-Assisting with Insertion and Removal. Policies and Procedures. Retrieved

from https://www.saskatoonhealthregion.ca/about/NursingManual/1186.pdf

Teleflex® Incorporated. (2015). ARROW®EZ-IO®: EZ-IO® Intraosseous Vascular Access Needles Instructions for Use. Retrieved June 6, 2018,

from https://www.teleflex.com/global/clinical-resources/ez-io/8082\_Rev\_02\_-FDA\_Intraosseous\_Infusion\_System\_IFU\_ATH\_v2\_- PRESS.pdf



## **HURON PERTH HEALTHCARE ALLIANCE**

## **Physician Approval Form**

<b>Medical Directive</b>	INTRAOSSEOUS INSERTION – PAEDIATRIC
Directive #	MD-ED-CC-028

## I, the undersigned physician, have:

- Reviewed the directive to fully understand the conditions under which it
  will be implemented, including knowing how the staff will be educated to
  provide this care and how they document or make me aware that the
  directive has been implemented so I can assume care appropriately, and
- Agree to assume the care of patients who have had an intervention performed as authorized by the directive

Name of Physician (please print)	
Signature	
Signature	
Date	

**Note:** The above Medical Directive was approved at the HPHA Medical Advisory Committee meeting November 22, 2018. If agreeable, please sign and return this form to Medical Services, Attn: Lori Hartman (Fax 519-271-7137).