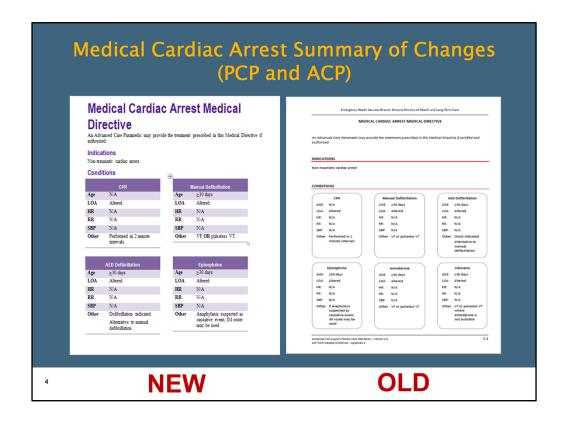


App Update: 4.0 will be updated on July 16<sup>th</sup>, 2017 – update your app (Ontario Paramedic Clinical Guide)

# Medical Directives Summary of Changes

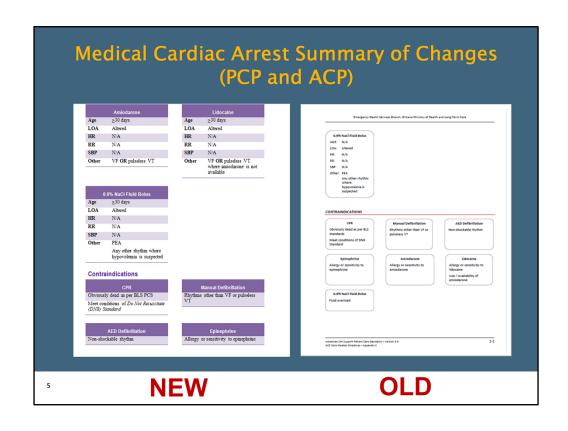
- Medical Cardiac Arrest
- ETCO2 (if applicable)
- Neonatal Resuscitation
- ROSC
- Cardiac Ischemia
- Hypoglycemia
- Mod-Severe Allergic Reaction
- Adrenal Crisis
- Tracheostomy Suctioning

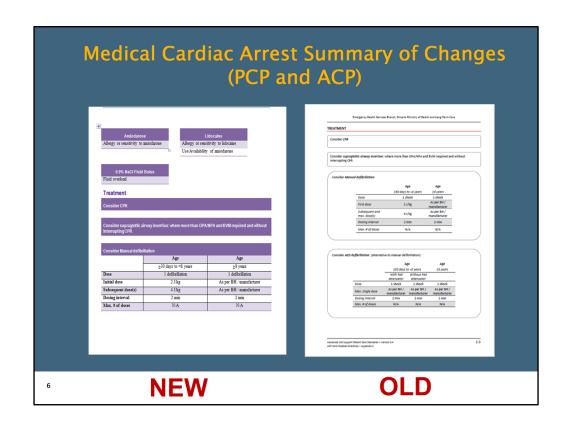
# MEDICAL CARDIAC ARREST

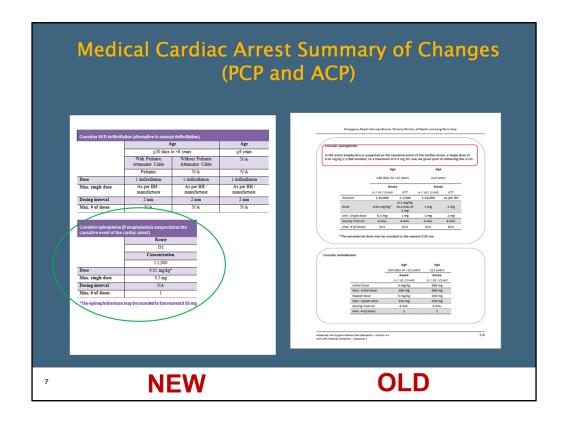


No change to the beginning of the cardiac arrest management plan Service specific teaching – SAED vs. Manual

Look of the directive has changed, but application hasn't changed





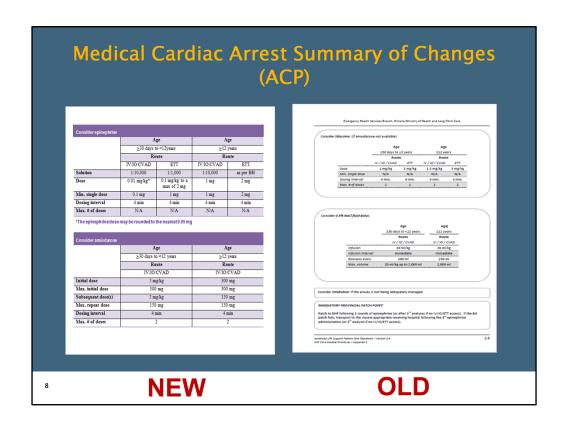


Anaphylaxis VSA – Consider Epi ASAP and early transport. Epi should not delay defibrillation.

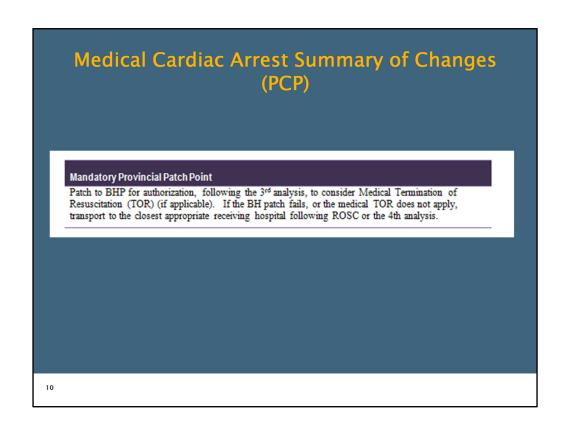
For ACPs in cases of VSA Anaphylaxis:

Following the first rhythm interpretation, if resources permit, establish an IV and administer Epi 1:10000 IV as indicated for cardiac arrest. If there is a delay to IV access, administer Epi 1:1000 IM.

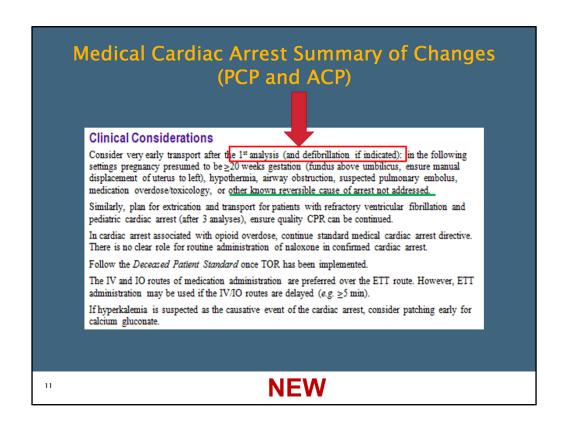
IFF Epi 1:10000 is administered IV, a subsequent dose of Epi 1:1000 IM is not required for anaphylaxis.



#### Medical Cardiac Arrest Summary of Changes (ACP) Age ≥30 days to <12 years Route IO/CVAD ETT ≥12 years Route IV/IO/CVAD Dose Min. single dose Dosing interval Max. # of doses 1.5 mg/kg N/A 4 min 3 mg/kg N/A 4 min 2 Consider 0.9% NaCl fluid bolus Age ≥12 years Route IV/IO/CVAD Route IV/IO/CVAD Infusion Infusion interval Reassess every Max. volume 20 ml/kg Immediate 100 ml 2,000 ml 20 ml/kg Immediate 250 ml 2,000 ml Patch to BHP following 3 rounds of epinephrine (or after 3<sup>rd</sup> analyses if no IV-IO ETT access). If the BH patch fails, transport to the closest appropriate receiving hospital following the 4<sup>rd</sup> epinephrine administration (or 4<sup>rd</sup> analysis in to IV/IO ETT access). **NEW**



If transport is required, consider appropriate airway management (King LT/ETT vs. OPA).



Key – early transport after first analysis – not first NSI

Email from Steph to Matt on the topic: Hey Matt,

The PHCSs and I are looking for clarification regarding the clinical considerations within the Medical Cardiac Arrest Medical Directive.

"Consider very early transport after the  $1^{st}$  analysis (and defibrillation if indicated) in the following settings: pregnancy presumed to be  $\geq$ 20 weeks gestation (fundus above umbilicus, ensure manual displacement of uterus to left), hypothermia, airway obstruction, suspected pulmonary embolus, medication overdose/toxicology, or other known reversible cause of arrest not addressed.

Similarly, plan for extrication and transport for patients with refractory ventricular fibrillation and pediatric cardiac arrest (after 3 analyses), ensure quality CPR can be continued."

Does the 2<sup>nd</sup> paragraph assume pediatric cardiac arrest particularly with refractory VF OR all pediatric arrests?

Prepare to transport regardless of rhythm, so all peds arrests. PCPs - Do not meet TOR criteria; ACPs - plan to transport to hospital because 99% of the time, will not get a field pronouncement

IFF the 2<sup>nd</sup> paragraph assumes pediatric cardiac arrest with refractory VF only (and not all peds arrests), then can non-refractory VF pediatric cardiac arrests be included in the 1<sup>st</sup> paragraph in the "or other known reversible cause" category?

We have interpreted that the majority of peds cases fall under reversible causes, so can transport after one analysis. However, the directive allows for use of clinical judgment, case by case(stay on scene for peds VF, vs leave after one analysis in presumed sepsis related peds VSA)

Rational: In the ALS PCS 3.4, we were to leave following the first NSI in the ped population. Now it appears we are to stay for 3.

No hard set rule based on our interpretation, given 1<sup>st</sup> paragraph allows for transport after one analysis in reversible causes (majority of peds arrests), or stay for 3 analysis prior to transport (ex patient in VF who may benefit from defibrillation).

#### The Airway Question

 In the ALS patient care standards it states that a Supraglottic Airway (King LT) is indicated when there is a "Need for ventilatory assistance OR airway control AND Other airway management is inadequate or ineffective."

#### Question:

 In the "un-controlled" world of EMS would it not be more effective to use a King LT instead of an oral airway after the first round of CPR is complete?
 Is it acceptable to use the King LT on VSA patients when the oral airway provides sufficient airway support?

. .

#### The Airway Question

- Answer:
  - · Use your educated clinical judgment

• Ask MAC: Date: 13-Mar-2012

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In wording for the use of the King LT/ETT, you are now able to use your educated clinical judgment to make the decision as to what method of airway management is best suited to you, your patient, and the situation. For example, if your patient initially meets the criteria for a Medical TOR (you think you're going down that pathway), and the OPA is providing adequate airway patency, then you are welcome to continue using it and not insert the King LT or ETT. Keep in mind, that if you're anticipating the transfer to the stretcher, trip down a flight of stairs, and the bumpy drive to the ED, then the King should be considered.

As for timing of King LT/ETT insertion; if you've decided that insertion is preferred or required, there is no hard and fast rule about when to do so. We have shared as a teaching point to make your first attempt following the first analysis as this allows both for pre-oxygenation (prior to the first analysis), and a second attempt should you need it (following the third analysis).

Bottom line: you're welcome to use either method, however, keep in mind the current and future management of the patient as the call progresses.

# END TIDAL CAPNOGRAPHY (ETCO2)

Confirm supraglotttic airway placeme	ent	
Method	Method	
Primary	Secondary	•
ETCO <sub>2</sub> (Waveform capnography)	ETCO <sub>2</sub> (Non-waveform device)	
	Auscultation	•
	Chest rise	
the mouth.  Confirmation of supraglottic airway n	ertion is defined as the insertion of the s must use ETCO <sub>2</sub> (Waveform capnograp working, then at least 2 secondary met	hy). If wavefor

Note: Biggest change is the NEED for primary confirmation for King LT/ETT

Services carrying ETCO2 must use it for the above.

#### ETCO2

Confirm orotracheal tube placement	
Method	Method
Primary	Secondary
ETCO <sub>2</sub> (Waveform capnography)	ETCO <sub>2</sub> (Non-waveform device)
	Visualization
	Auscultation
	Chest rise
	Esophageal detection device

#### **Clinical Considerations**

An intubation attempt is defined as insertion of the laryngoscope blade into the mouth for the purposes of intubation.

Confirmation of orotracheal intubation must use ETCO<sub>2</sub> (Waveform capnography). If waveform capnography is not available or not working then at least 3 secondary methods must be used. Additional secondary ETT placement confirmation devices may be authorized by the local medical

ETT placement must be reconfirmed immediately after every patient movement.

## ETCO2 Changes Review (ACP only)

Confirm nasotracheal tube placement	1
Method	Method
Primary	Secondary
ETCO <sub>2</sub> (Waveform capnography)	ETCO <sub>2</sub> (Non-waveform device)
	Auscultation
	Esophageal detection device
	Chest rise

#### **Clinical Considerations**

A nasotracheal intubation attempt is defined as insertion of the nasotracheal tube into a nare.

Confirmation of nasotracheal placement must use  ${\rm ETCO_2}$  (Waveform capnography). If wave-form capnography not available or not working, then at least 2 secondary methods must be used ETT placement must be reconfirmed immediately after every patient movement.

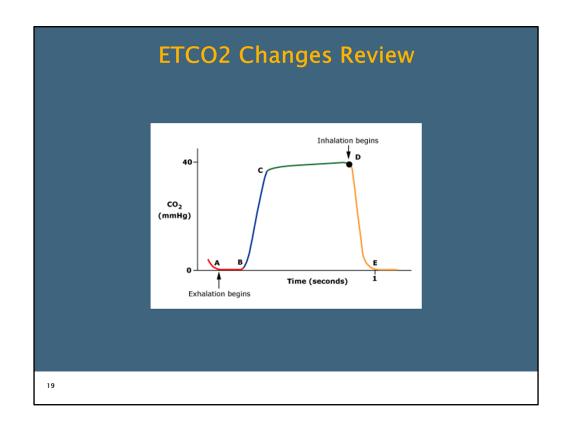
## ETCO2 Changes Review (ACP only Service dependent)

Consider cricothyrotomy tube placen	nent
Method	Method
Primary	Secondary
ETCO <sub>2</sub> (Waveform capnography)	ETCO <sub>2</sub> (Non-waveform device)
	Auscultation
	Chest rise

#### **Clinical Considerations**

Confirmation of cricothyrotomy must use ETCO<sub>2</sub> (Waveform capnography). If waveform capnography is not available or not working, then at least 2 secondary methods must be used. Additional secondary Cricothyrotomy tube placement confirmation devices may be authorized by the local medical director.

Cricothyrotomy tube placement must be reconfirmed immediately after every patient movement.



Describe the inhalation/exhalation cycle

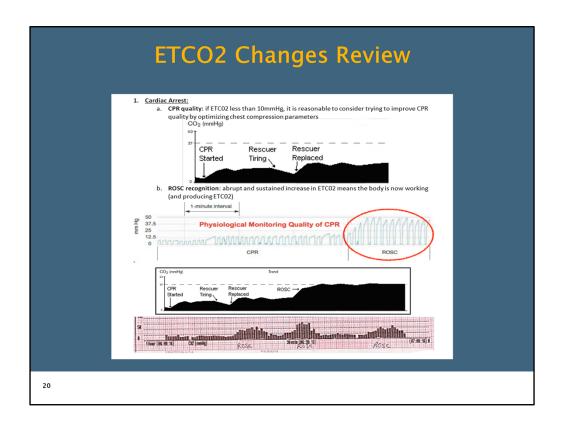
Phase A is the start of the respiratory baseline

Phase B is the end of the respiratory baseline and the exhalation begins

Phase C is the expiratory plateau (which means the end of the breathing out)

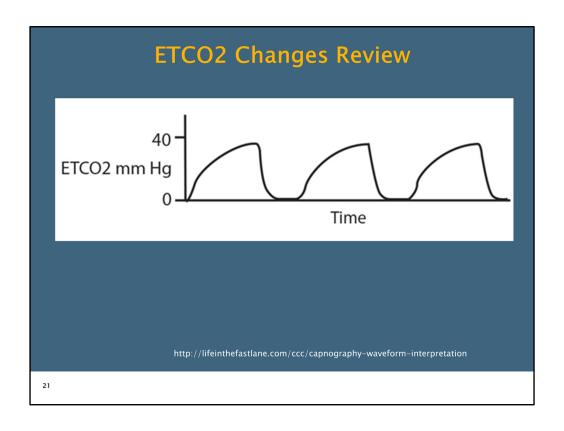
Phase D is the end of the exhalation phase and this is where the ETCO2 value or number is measured from. The highest concentration of CO2 is at this point.

Phase E is when the patient takes a breath in and there is a drop in the CO2 as there is nothing being exhaled.



Uses of ETCO2 in the VSA pt

- -positioning of ETT
- -quality of CPR
- -possible indicator of ROSC (with spike in ETCO2)
- -possible factor associated with determining/granting TOR



Shark fin – evidence of bronchoconstriction

#### In regards to the Shark Fin presentation:

Phases A and B remain essentially unchanged.

Phase C has a more pronounced slope during the expiratory phase and this is caused from the air trapping and bronchospasm that is taking place which thereby inhibits CO2 from being released more efficiently hence when they say that COPD'ers are retainers; that's why they are!

Phase D remains unchanged as well.

Phase E is when the patient takes a breath in and there is a drop in the CO2 as there is nothing being exhaled.

#### ETCO2

In the DKA patient ETCO2 varies directly with the patient's pH. So the farther out of usual range the ETCO2 is, the sicker the patient.

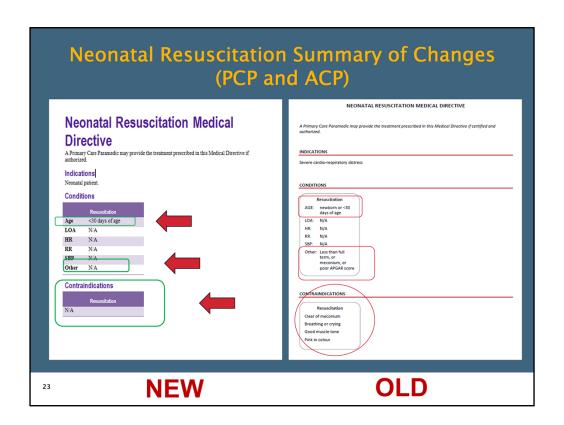
#### In regards to Early recognition of Sepsis:

The initial stage of SIRS deals with vital signs assessment:

Heart rate greater than 90
Breathing rate greater than 20
Temp greater than 38 or less than 36
ETCO@ less than 32

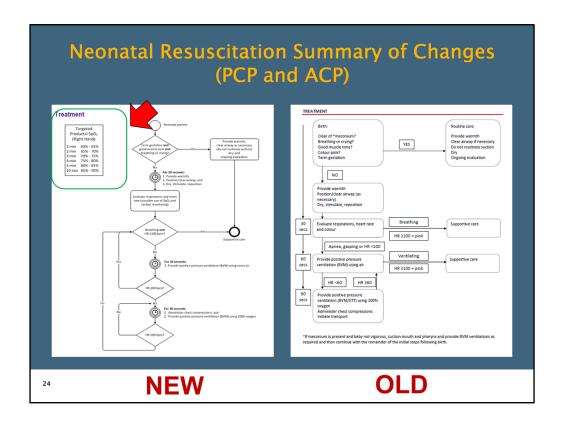
Sick patients breath faster and blow off more CO2; therefore less than normal values of 35-35 mmHg TORR. So the sick patient in the nursing home with a UTI on an antibiotic and tachycardia, with a low end tidal means sick patient. All patients can be monitored for their CO2 and should be.

# NEONATAL RESUSCITATION



#### Wording change

- Indications
- Conditions
- Contraindications removed



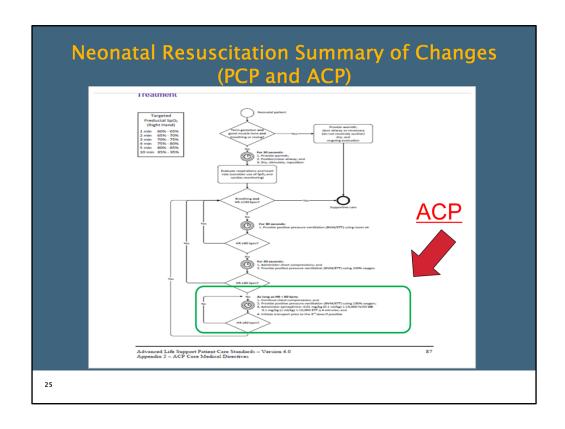
Wording changes: No longer evaluate colour

Algorhythm format change

Same treatment with the addition of cardiac and sp02 monitoring – only applicable to newborn (JUST been born).

Newly born – start at the top

Neonate – start at 2<sup>nd</sup> box (skip the first 30s)



Main Focus is to initiate transport prior to the 3<sup>rd</sup> dose if possible

# Neonatal Resuscitation Summary of Changes (PCP and ACP)

#### **Clinical Considerations**

If neonatal resuscitation is required, initiate cardiac monitoring and pulse oximetry monitoring.

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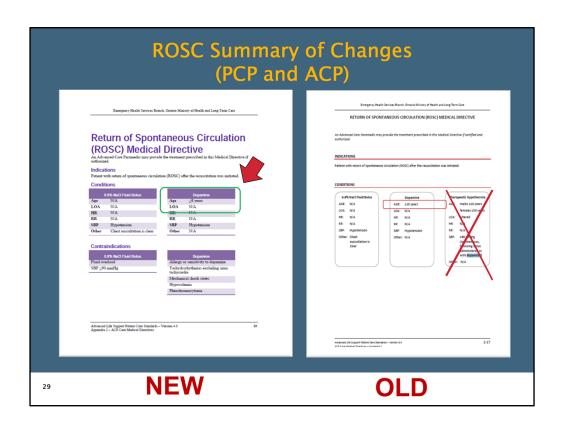
**NEW** 

#### Neonatal Resuscitation Recap

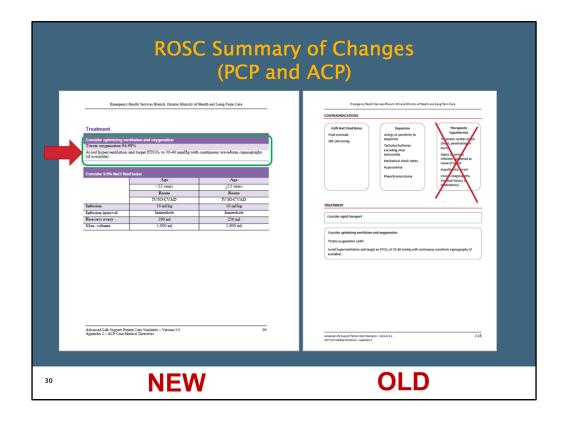
- Less 'wordy'
  - i.e. indications: Neonatal (<30d) patient
- Algorithm format change
- Same assessment with the exception of:
  - Evaluation of colour
- Addition of cardiac monitor and Sp02 during resuscitation



### RETURN OF SPONTANEOUS CIRCULATION (ROSC)

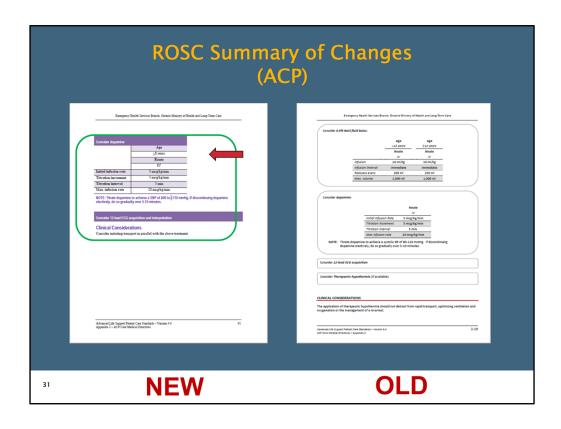


Dopamine ≥8y



NEW – target SPO2/ETCO2

Normal ETCO2 is 35-45, but our target is 30-40 (hyperventilation)



Note: Single strength dopamine

## ROSC Summary of Changes (ACP)

- Question: Within the ROSC Medical Directive, the ONLY route identified for the administration of a fluid dopamine is via IV. Can we administer dopamine via IO or CVAD?
- Answer: Maybe...
- Ask MAC: Date: <u>15-Feb-2012</u>

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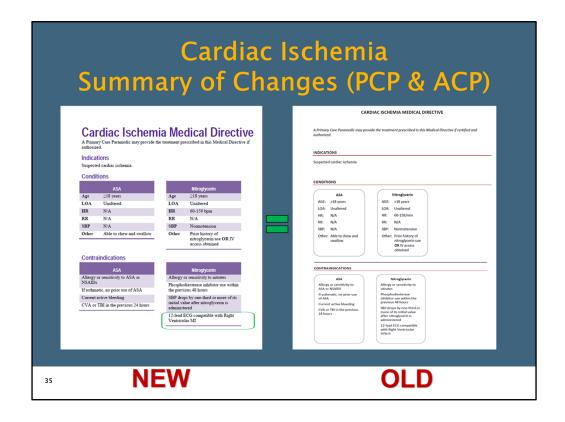
#### Answer:

IO and CVAD are not identified as routes of administration for dopamine within the medical directive. There is concern that extravasation may lead to tissue destruction, however the same concern exists for IV administration. For this reason, should you feel administration of dopamine via IO or CVAD is appropriate for the patient, a patch to the BHP is required.

		(ACI	1		4	
Single Stre	ngth Dopamino DOPAMINE INFUSI (Using a		or drops/min with	n a microdrip set)		
		Drij	p Rate (drops/mir	ate (drops/min)		
Weight (kg)	(mcg/kg/minute)	5 (mcg/kg/minute)	10 (mcg/kg/minute)	15 (mcg/kg/minute)	20 (mcg/kg/minute)	
5	1	2	4	6	8	
10	2	4	8	11	15	
15	3	6	11	17	23	
20	4	8	15 19	23	30 38	
30	5	11	23	34	45	
35	5	13	26	39	53	
40	6	15	30	45	60	
45	7	17	34	51	68	
50	8	19	38	56	75	
55	8	21	41	62	83	
60	9	23	45	68	90	
65	10	24	49	73	98	
70	11	26	53	79	105	
75 80	11	28 30	56 60	90	113 120	
85	13	32	64	96	128	
90	14	34	68	101	135	
95	14	36	71	107	143	
100	15	38	75	113	150	
105	16	39	79	118	158	
110	17	41	83	124	165	
115	17	43	86	129	173	
120	18	45	90	135	180	

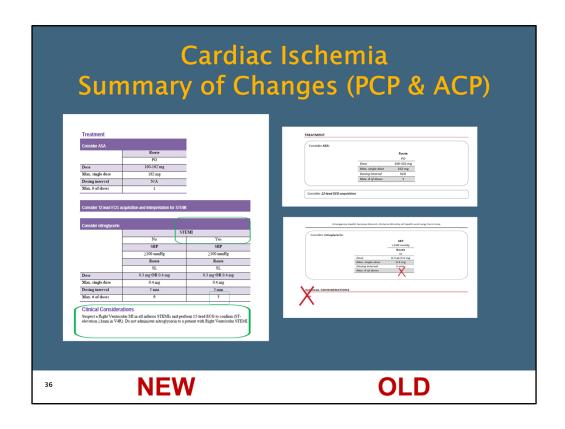
As per the equipment standards this is changing to 800mcg,

# Cardiac Ischemia



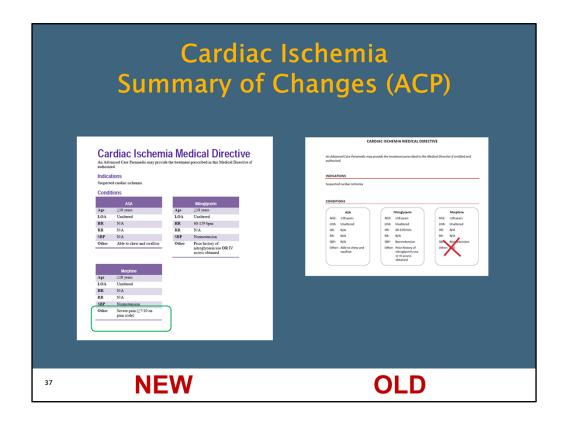
Indication, Conditions and Contraindications: wording remains the same.

However, now medics will be REQUIRED to rule out a RVMI prior to administering NTG.



Yes – start performing a modified 12 Lead

Max 3 NTG for STEMI+



#### Rational for LESS morphine:

#### Ischemic CP

Morphine was previously a class I recommendation, but is now bumped down to class II (conflicting evidence). Therefore decision made to be more selective with who gets morphine.

MAC Summary: Platelets inhibition is an integral part of STEMI therapy and highrisk ACS patients. Suboptimal platelet inhibition early after primary stent implantation for STEMI is associated with thrombotic complications, including stent thrombosis. (13) Thus, given the evidence of reduced platelet inhibition with concomitant morphine administration, and in the absence of data that may support a potential clinical benefit of morphine in ACS, more caution should be used regarding morphine administration, and a selective rather than routine morphine use seems to be reasonable. Morphine administration might be reserved for ACS patients presenting with persistent severe chest pain. Severe chest pain has been previously defined as a self- reported numeric rating scale ≥ 7, based on a scale from 0 to 10 as described in cancer patients treated with Morphine.

MAC Recommendation: Morphine use in ACS patients be restricted to those with severe pain ( $\geq$  7), after other anti-ischemic therapies have been started.

#### Cardiac Ischemia Summary of Changes (PCP and ACP)

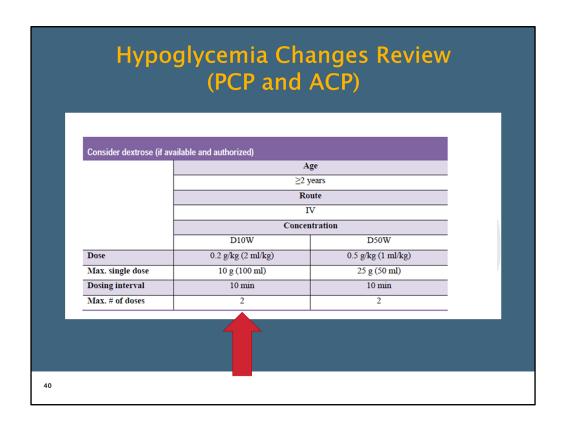
- A 68 year old male is complaining of ischemic-type chest discomfort. He has been instructed by the 911 call taker to take his own ASA.
  - •Do you administer more?

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Ontario Central Ambulance Communications Centres have begun directing patients with chest pain, who already take aminosalicylic acid (ASA) to chew their own ASA (160-325 mg). ASA has demonstrated benefit to outcomes in patient suffering myocardial ischemia or infarction. As ASA takes time to be absorbed, dispatch-directed early administration has potential benefits, is recommended as the standard of care in the Heart and Stroke Foundation of Ontario Advanced Cardiac Life Support Guidelines 2015, and poses minimal risk to the patients.

ASA is a safe medication with a wide therapeutic index (the effective dose without side effects can be from 80 to 1500 mg). The additional dose provided by paramedics will not exceed the therapeutic dose, and there is a chance the patient did not have or did not take the correct dose suggested by the ambulance communications officer. Therefore, paramedics should continue to follow their Ischemic Chest Pain Medical Directive without modification when patients have already taken ASA prior to paramedic arrival.

# Hypoglycemia



Should service carry D10W; otherwise no change

NO titration to effect with D10 or D50

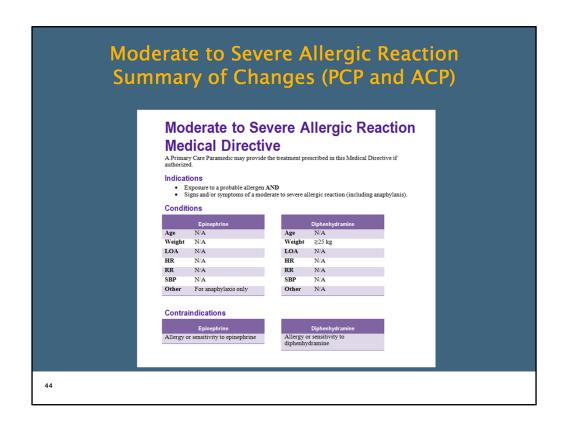
### Hypoglycemia Changes Review (ACP only)

Consider dextrose (D10)	W pre-mixed)		
	Age	Age	
	<30 days	≥30 days	
	Concentration	Concentration	
	D10W	D10W	
	Route	Route	
	IV	IV	
Dose	0.2 g/kg (2 ml/kg)	0.2 g/kg (2ml/kg)	
Max. single dose	5 g (50 ml)	10g (100 ml)	
Dosing interval	10 min	10 min	
Max. # of doses	2	2	

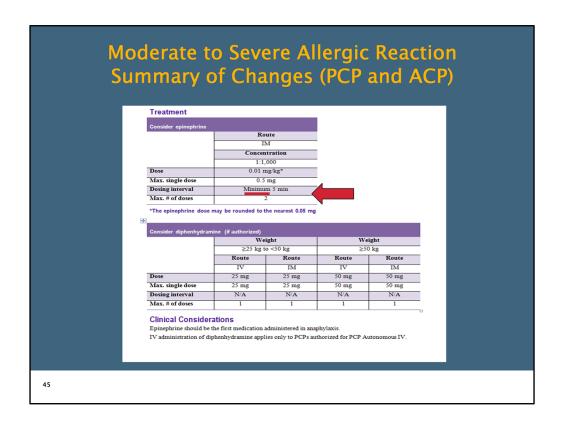
### Hypoglycemia Changes Review (ACP only)

Consider dextrose (D50)	W diluted as required if no	ot using D10W)	
	Age	Age	Age
	<30 days	≥30 days to <2 years	≥2 years
	Concentration	Concentration	Concentration
	D10W	D25W	D50W
	Route	Route	Route
	IV	IV	IV
Dose	0.2 g/kg (2 ml/kg)	0.5 g/kg (2 ml/kg)	0.5 g/kg (1 ml/kg)
Max. single dose	5 g (50 ml)	10 g (40 ml)	25 g (50 ml)
Dosing interval	10 min	10 min	10 min
Max. # of doses	2	2	2

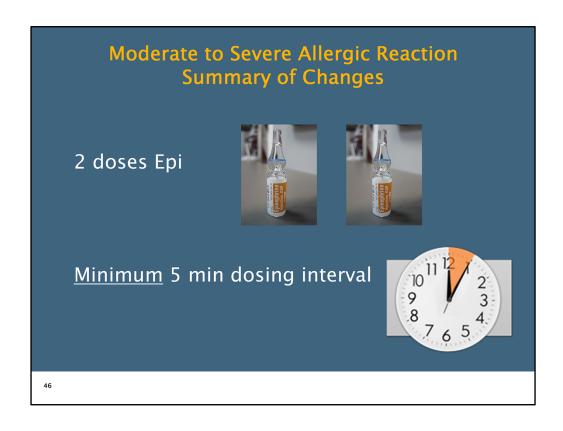
## MODERATE TO SEVERE ALLERGIC REACTION



No changes

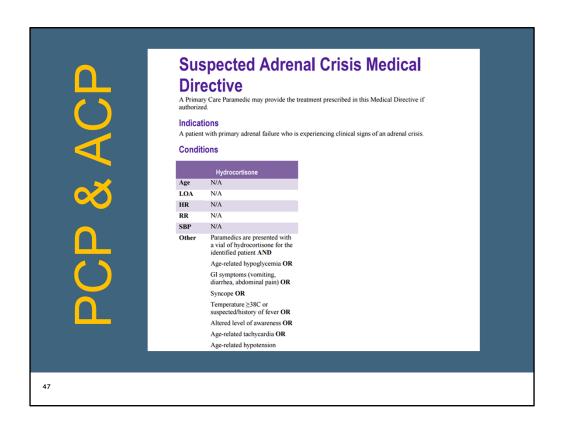


Change: max # doses 2; minimum 5 minute interval



When appropriate, follow up with Benadryl

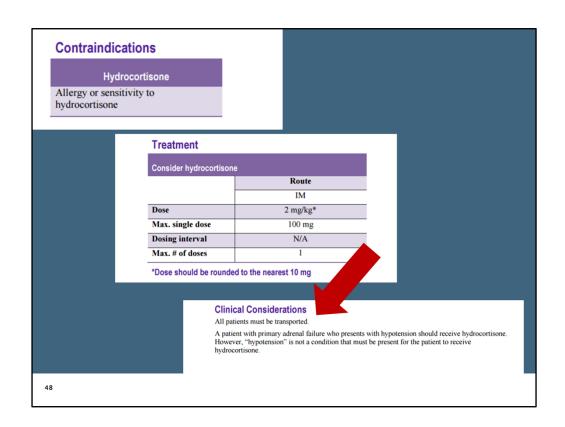
Do not give Benadryl and Gravol together



PCP and ACP same directive?

What's the "give early" rationale all about?

Early = less crisis-y (main goal to decrease hypotension)



What's the dosing?

Why "must" patients be transported? – significant follow up required in ED What do you do in hospital for these folks?



That's a hyperlink to 51 sec video

How do we give it?
Onset, effects and duration?

# ENDOTRACHEAL AND TRACHEOSTOMY SUCTIONING

Suctioning Medical Directive  A Primary Care Paramedic may provide the treatment prescribed in this Medical Directive if authorized.  Indications
Patient with endotracheal or tracheostomy tube;
AND
Airway obstruction or increased secretions.
Conditions
Suctioning
Age N/A
LOA N/A
HR N/A
RR N/A
SBP N/A
Other N/A
Contraindications
Suctioning

NEW

Important = PPE (mask/goggles/splash guard/gown)

Treatment			
Consider suctioning			
	Infant	Child	Adult
Dose	suction at 60-100 mmHg	suction at 100-120 mmHg	suction at 100-150 mmHg
Max. single dose	N/A	N/A	N/A
Dosing interval	1 minute	1 minute	1 minute
Max. # of doses	5	5	5
Clinical Consid Pre-oxygenate with In an alert patient, w Do not exceed 10 sec	100% oxygen. henever possible, have patie	ent cough to clear airway pr	ior to suctioning.

Start suctioning pressures low and increase as needed

NEW – max 5 # of doses

Reminder: Ventilate/oxygenate during the 1 min interval between suctioning attempts

