

Obstetric Management of Postpartum Hemorrhage

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Defining PPH

- Primary vs. Secondary
 - Primary
 - First 24 hours after delivery
 - Secondary
 - late or delayed PPH
 - between 24 hours and 12 weeks after delivery
- Classically defined by volume of blood loss
 - ≥ 500 ml at vaginal delivery
 - ≥ 1000 ml at cesarean delivery



Defining PPH cont.

- Royal College
 - Minor 500 1000 ml
 - Major > 1000 ml
 - moderate 1000 2000 ml
 - severe > 2000 ml
- International Expert Panel
 - "active bleeding > 1000 ml within the 24 hours following birth that continues despite the use of initial measures, including first-line uterotonic agents and uterine massage"



Incidence

- Varies widely depending on the criteria used to define
- US Data suggests incidence of 2-3% of deliveries (range most quoted is 1-5%)



Causes

- Atony
 - most common etiology
 - -~80% of cases
 - diagnosis made clinically
- Lacerations
 - cervical or vaginal lacerations
 - extensions of hysterotomy
 - ruptured uterus
- Retained products of conception
- Coagulopathy / medical bleeding



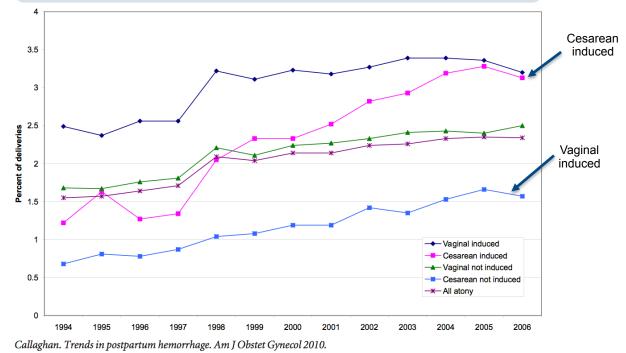


ATONY



Incidence of atony with PPH

Annual rates of postpartum hemorrhage caused by atony, by mode of delivery, and by induction status (United States, 1994–2006)







Incidence of atony with PPH

Delivery type	Rate (95% CI)			Odds ratio (95% CI)	
	1994	2006	Change, %	1994	2006
Overall	1.6 (1.4–1.8)	2.4 (2.2–2.5)	50		
Vaginal					
Induced	2.5 (2.3–2.8)	3.2 (3.0–3.5)	28	3.7 (3.2–4.4)	2.1 (1.9–2.2)
Not induced	1.7 (1.5–1.9)	2.5 (2.3–2.7)	47	2.5 (2.2–2.9)	1.6 (1.5–1.7)
Cesarean					
Induced	1.2 (0.9–1.6)	3.1 (2.8–3.5)	160	1.8 (1.4–2.3)	2.0 (1.8–2.2
Not induced	0.7 (0.6–0.8)	1.6 (1.5–1.8)	130	Reference	Reference





Uterine atony pathogenesis

- Characterized by inadequate uterine contraction following delivery
 - uteroplacental arteries remain open
 - local hemostatic factors unable to contribute
- Etiology not known
 - OXTR desensitization as a possible explanation
- Well-described risk factors
 - but not all present with risk factors





Prevention?



Prevention of uterine atony

- Active vs. expectant management of the third stage of labor
 - Active management
 - prophylactic uterotonic
 - early cord clamping
 - controlled cord traction
 - Expectant management
 - await signs of placental separation
- Pluses and minuses to both approaches
 - Utilize prophylactic uterotonics almost universal





Treatment

- Make the diagnosis of PPH due to atony
 - highly subjective
 - with experience..."that's too much"
 - tools exist to estimate blood loss
 - graduated measurement containers and drapes
 - · visual aids that provide quantified examples
 - wet-dry weights
 - newer optical technologies
 - early intervention
 - prevent shock, coagulopathy
 - rule out lacerations and retained products



- Unit-based PPH Protocols
 - allow for standardized management
 - multidisciplinary team approach
 - improved outcomes
 - time to cessation of bleeding
 - rates of DIC
 - decreased use of fewer blood products
 - California Maternal Quality Care Collaborative OB Hemorrhage Protocol





- Medical management
 - first line
 - utilize various uterotonic agents
 - each target different receptors or signaling pathways
- Blood component therapy
- Surgical management



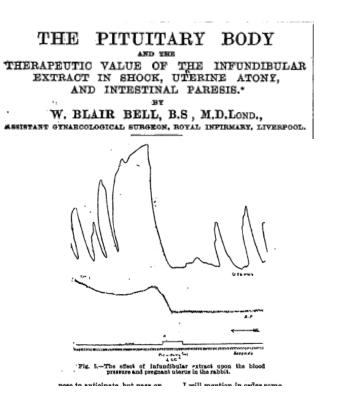


Oxytocin



Oxytocin history

- W. Blair Bell, 1909
 - British obstetrician
 - First to publish the clinical application of posterior pituitary extract, 1909
 - Administered IM
 - Prophylaxis and treatment of postpartum hemorrhage
 - First to use in Obstetrics
 - "There is no doubt, I think, that any one of these properties is sufficient to secure a permanent place in therapeutics for this much neglected portion of our anatomy."
 - "it ought but rarely to be given before delivery."
 - Concerned about tetanic contraction effects on fetus







Oxytocin history cont.

THE SEQUENCE OF AMINO ACIDS IN OXYTOCIN, WITH A PROPOSAL FOR THE STRUCTURE OF OXYTOCIN*

By VINCENT du VIGNEAUD, CHARLOTTE RESSLER, and STUART TRIPPETT

(From the Department of Biochemistry, Cornell University Medical College, New York, New York)

(Received for publication, July 13, 1953)

- du Vigneaud, 1953
 - In 1949, isolated pure oxytocin from the posterior pituitary
 - In 1953, proposed the amnio acid sequence and structure for oxytocin
 - Produced a synthetic product indistinguishable from the natural hormone
 - Won the Nobel prize for this discovery in 1955





- Oxytocin cont.
 - uterotonic agent of choice
 - no absolute contraindication
 - utilize even after prophylactic usage
 - various regimens
 - 40 Units in 1 liter of normal saline or LR
 - deliver via pump
 - rate to control hemorrhage
 - · adjust rate to achieve/maintain contractility
 - 10 units IM if no IV access
 - avoid undiluted IV boluses

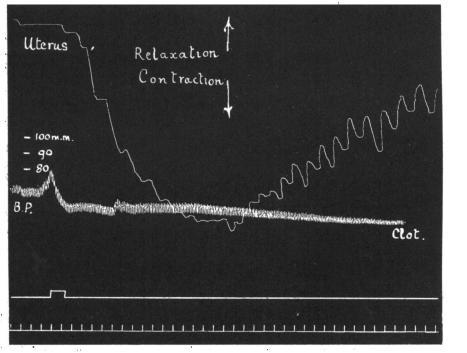




Ergots

ON SOME PHYSIOLOGICAL ACTIONS OF ERGOT. By H. H. DALE.

(From the Wellcome Physiological Research Laboratories, Herne Hill.)





Dale. 1906



- Ergots cont.
 - methylergonovine 200 mcg IM
 - do not give IV
 - avoid in
 - hypertension, Raynaud's
 - response in 2-5 minus
 - if first dose ineffective, switch to next agent
 - dosing intervals every 2-4 hours up to 1mg





- Carboprost
 - PGF₂alpha (Hemabate) 250 mcg IM
 - do not give IV
 - avoid in asthma/bronchospasm
 - dosing intervals every 15-90 mins up to 8 doses, 2 mg
 - side effects
 - tachycardia, fever, diarrhea





- Misoprostol (PGE1)
 - useful when other uterotonic agents are contraindicated
 - optimal dose and route unclear
 - sublingual
 - 400 mcg
 - rapid absorption, 30 min peak,
 - avoids first-pass (higher peak, longer affect)
 - oral
 - 400 mcg
 - rapid absorption, 30 min peak
 - first pass effect (lower peak, shorter affect)
 - rectal
 - 800-1000 mcg
 - slow onset (one hour) but long duration (four hours)





- Misoprostol (PGE1) cont.
 - fever common
 - not contraindicated in hypertension or asthma



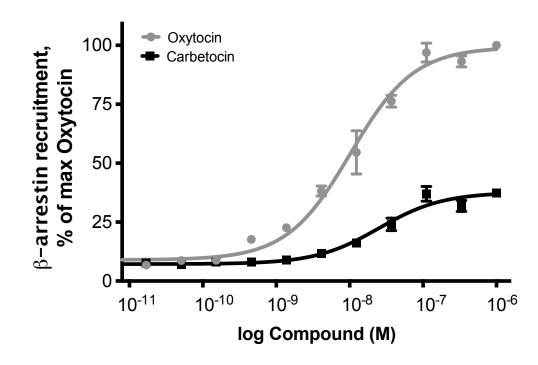


- Carbetocin
 - long-acting oxytocin analog
 - modified to have a long serum $^{1\!\!/_2\text{--life}}$
 - not available in US
 - primarily used for prevention of PPH following cesarean delivery
 - marketed due to it's long ½-life, therefore longer duration of action...but





• Carbetocin cont.







- Tranexamic acid
 - anti-fibrinolytic agent
 - used for prevention and treatment of PPH
 - WOMAN trial on-going
 - early use of TXA in women with PPH
 - primary outcome
 - mortality, hysterectomy, severe morbidity





- Explore uterine cavity
 - evacuate clots
 - facilitates further uterine contraction





Duke L&D Bedside Cognitive Aid

Drug	Dose	Route	Frequency	Contraindications
Oxytocin — 1 st Line	10-40 Units per 500-1000 ml	IV Infusion (10 Units IM)	Continuous infusion	Avoid undiluted rapid infusion
Methylergonovine (Methergine)	0.2 mg	IM (NOT IV)	Every 2-4 hours	Avoid if hypertensive
Prostaglandin F2α (Hemabate)	250 mcg	IM (NOT IV)	Every 15-90 mins (limit 8 doses in 24 hrs.)	Avoid in asthmatic
Misoprostol (Cytotec, PGE1)	800-1000 mcg	Rectal	Once	Rare
Tranexamic acid* (TXA)	1 g bolus	IV	May follow with infusion (1mg/kg/hr)	Avoid if hypertensive or recent DVT/PE

*Consult with Ob Anesthesia prior to giving





Uterine atony surgical treatment

- Surgical treatment
 - Occlude uterine cavity
 - Bakri balloon
 - Uterine packing
 - B-Lynch uterine suture braces
 - Uterine devascularization
 - Uterine artery ligation
 - Utero-ovarian vessel occlusion
 - Uterine artery embolization
 - Hysterectomy





• Uterine Balloons

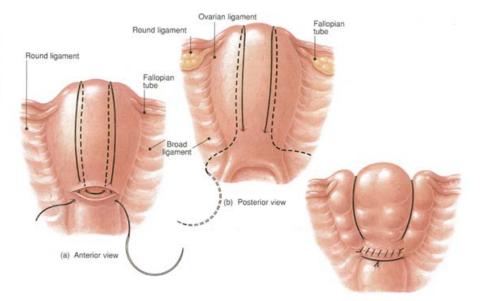




https://www.cookmedical.com/products/wh_sosr_webds/



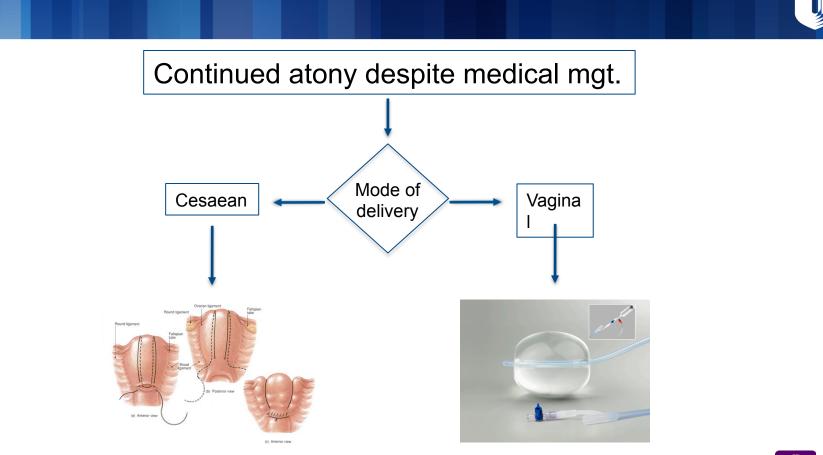
• B-Lynch Uterine Suture Brace







http://www.cblynch.co.uk/wp-content/uploads/tech-image.jpg





Lacerations

- Cervical and Vaginal lacerations
 - identified on initial presentation with PPH
 - repair



Retained products

- Careful visualization of placenta
- Manual uterine exploration
- Ultrasound



Fluids and blood products

• Discussed separately





Obstetric management of PPH

- Summary
 - Preparation and anticipation
 - Early diagnosis
 - Rule out lacerations and retained products
 - Uterine atony Rx
 - medical mgt
 - more medical mgt
 - more medical mgt
 - surgical options
- Research gaps
 - optimal drug combinations
 - personalized uterotonic approach to treatment





Questions?