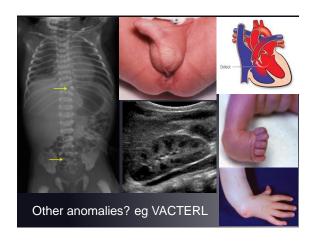
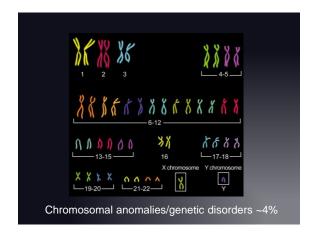


Taghavi & Stringer 2017





Incidence: 1:3-4000 live births so ~18/yr in NZ

Aetiology: sporadic (MZ twins 2.5% concordance), recurrence 1%

Other anomalies >50% e.g. VACTERL, cong heart disease

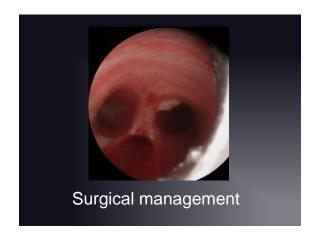
Antenatal clues: polyhydramnios & absent/small stomach

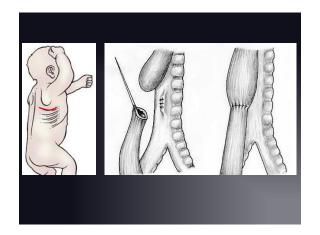
Postnatal: mucousy baby who chokes/ coughs when fed

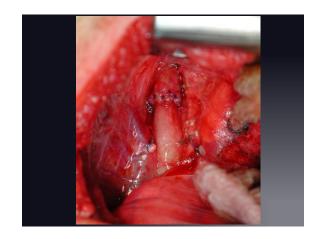
Diagnosis: pass a 10Fg gastric tube and take CXR + AXR



Nurse head up ± semi prone to keep upper pouch empty
 Transfer to NN surgical unit promptly (urgently if respiratory distress or needing ventilation)
 Invs: echo, X-rays, renal US etc
 Non urgent 1º repair if stable







Uneventful recovery in many – extubation, temporary tube feeding and then oral feeding

Survival (Spitz classification)

Birth wt.>1500g and no major cardiac anomaly 98%

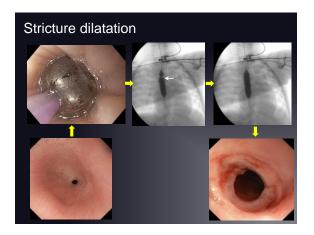
Birth wt. <1500g or major cardiac anomaly 82%

Birth wt. <1500g plus major cardiac anomaly 50%

.....and improving

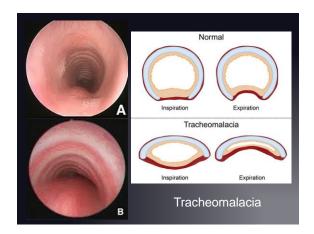


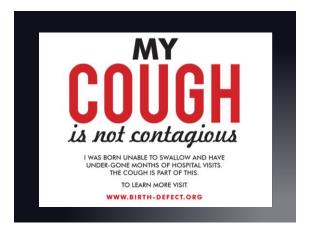


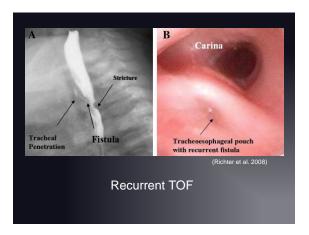






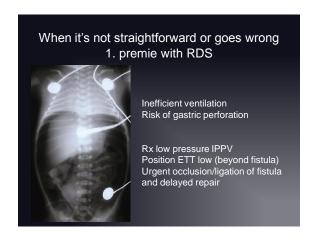


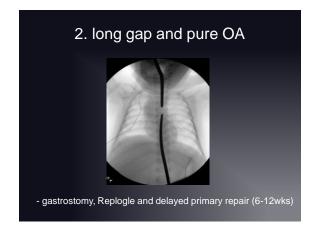






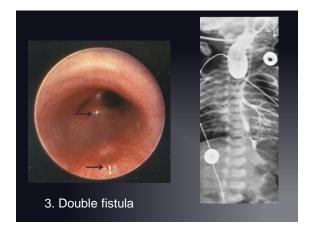


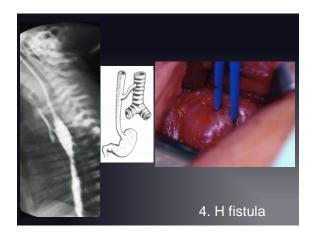




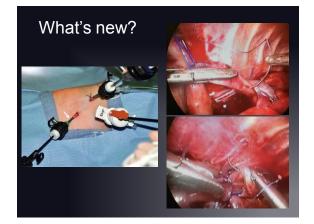












Key messages

- Consider OA in any newborn with maternal polyhydramnios
- If OA suspected try and pass a 10FG gastric tube and take an X-ray if it won't enter stomach
- OA/TOF + respiratory distress = urgent transfer
- Complications after OA repair (leak, stricture, GOR, dysmotility) are manageable
- With appropriate nurture, outcomes are generally good but more guarded if major CHD