

BEYOND SIMPLE GASTRO

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CASE PRESENTATION

A 2 day old female infant presents with profuse watery diarrhoea.

Investigations:

Na 130mmol/l

K 3.5mmol/l

Faecal osmolar gap 120

Stool MCS - negative

INVESTIGATIONS

FAECAL OSMOLAR GAP

Serum osmolarity – $2 \times (\text{Sodium} + \text{Potassium faecal conc})$

$<50 \text{ mosmo/l}$ = secretory diarrhoea

$50\text{-}100 \text{ mosmo/l}$ = could be either

$>100 \text{ mosmo/l}$ = osmotic diarrhoea

Need a liquid stool

FAECAL REDUCING SUBSTANCES

Positive = osmotic diarrhoea

DIFFERENTIAL DIAGNOSIS

- * Congenital enteropathy
- * Cow's/soya milk protein allergy
- * Cystic Fibrosis
- * Lactose intolerance
- * Congenital chloride diarrhoea
- * Congenital sodium diarrhoea

CONGENITAL ENTEROPATHY

- * Diarrhoea shortly after birth
- * TPN dependant

- * Congenital microvillus inclusion disease
- * Congenital tuft enteropathy
- * Treatment – life long TPN or small bowel transplant

COW'S MILK PROTEIN ALLERGY

- * No gold standard investigation
- * Eliminate cow's milk from the diet
- * A 2 week trial of amino acid base formula
- * If symptoms resolve, re-introduce cow's milk
- * If symptoms recur = Cow's milk protein allergy

CYSTIC FIBROSIS

- * Most common autosomal recessive genetic disorder
- * Incidence
- * Presents with bulky, smelly frequent stools
- * Severe failure to thrive or severe protein energy malnutrition with oedema
- * Stool for free fat
- * Faecal elastase - $<15\mu\text{g/g}$ - $<200\mu\text{g/g}$
- * Gold standard – Sweat test – $\text{Cl} >60$

CONGENITAL CHLORIDE DIARRHOEA

- * Primary defect in the active transport of Cl/HCO_3 in the distal ileum and colon
- * Onset is intrauterine, leading to watery diarrhoea and polyhydramnios.
- * There is no passage of meconium due to watery, 'urine like' diarrhoea.
- * Watery diarrhoea may be easily confused with urine.
- * Hypochloraemia, hyponatraemia, hypokalaemia and metabolic alkalosis

CONGENITAL CHLORIDE DIARRHOEA

- * CLD is then confirmed by faecal electrolytes.
- * A faecal chloride concentration exceeding the sum of the sodium and potassium concentrations suggests the diagnosis of CLD.
- * CLD must be considered in the differential diagnosis in children with a tendency to chronic dehydration, failure to thrive, slow growth, and hypokalaemic and hypochloraemic metabolic alkalosis.

CONGENITAL SODIUM DIARRHOEA

- * The diagnosis based on
 - daily faecal sodium excretion between 98 and 190 mmol/L
 - hyponatremia,
 - metabolic acidosis
 - low-to-normal urinary sodium concentrations

CONGENITAL SODIUM DIARRHOEA

- * Inborn defect of intestinal sodium/proton exchange
- * Severe diarrhoea with voluminous alkaline stools
- * Contains high concentrations of sodium and resulting in hyponatremia and metabolic acidosis.
- * Management is with total parenteral nutrition

LACTOSE INTOLERANCE

- * Primary is rare
- * Most cases secondary post-enteritis
- * Diagnosis- reducing substances positive in the stool
- * Eliminate lactose

COELIAC DISEASE

- * Symptoms start on introduction of gluten
- * Loose stools, abdominal bloating, severe microcytic anaemia, oedema and failure to thrive
- * High anti-tissue transglutaminase, anti-endomysial antibodies
- * Gold standard- villous atrophy on duodenal biopsy
- * Eliminate gluten from the diet

TODDLER'S DIARRHOEA

- * Presents with loose stools with undigested food visible
- * Not dehydrating
- * No failure to thrive
- * Might be aggravated by large volumes of fruit juice
- * No treatment needed, reassure

IRRITABLE BOWEL SYNDROME

- * Older child
- * Stools range from loose to hard
- * Associated with abdominal pain
- * Relieved by stooling
- * No failure to thrive
- * Can attempt dietary changes
- * Symptomatic relief

INFLAMMATORY BOWEL DISEASE

- * Mucoid bloody stools
- * Recurrent abdominal pain
- * Loss of weight
- * Painless anal fissures
- * Mouth ulcers
- * Fever

INFLAMMATORY BOWEL DISEASE

- * Faecal calprotectin
- * By product of neutrophil breakdown on the mucosal surface
- * A marker of intestinal inflammation
- * Level of 200ug/g significant
- * Screening for colonoscopy

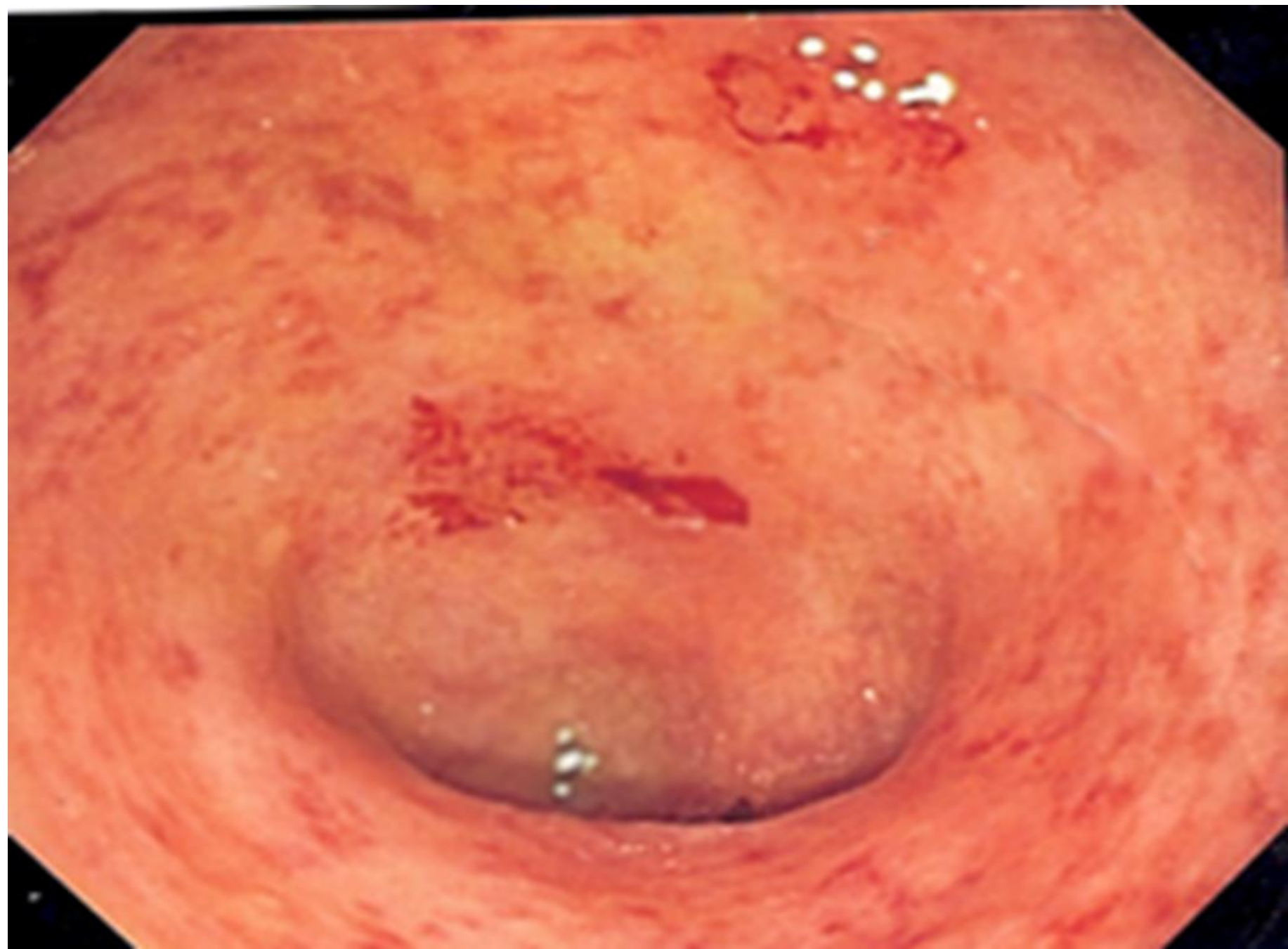
CROHN'S DISEASE

- * Affects mouth to anus
- * Causes strictures and fistulae
- * Mostly affects the terminal ileum
- * 30% small bowel involvement, 30% colon only involvement, 40% small and colon
- * Mild disease – Enteral nutrition
- * Moderate to severe disease – Steroids, azathioprine or monoclonal antibodies



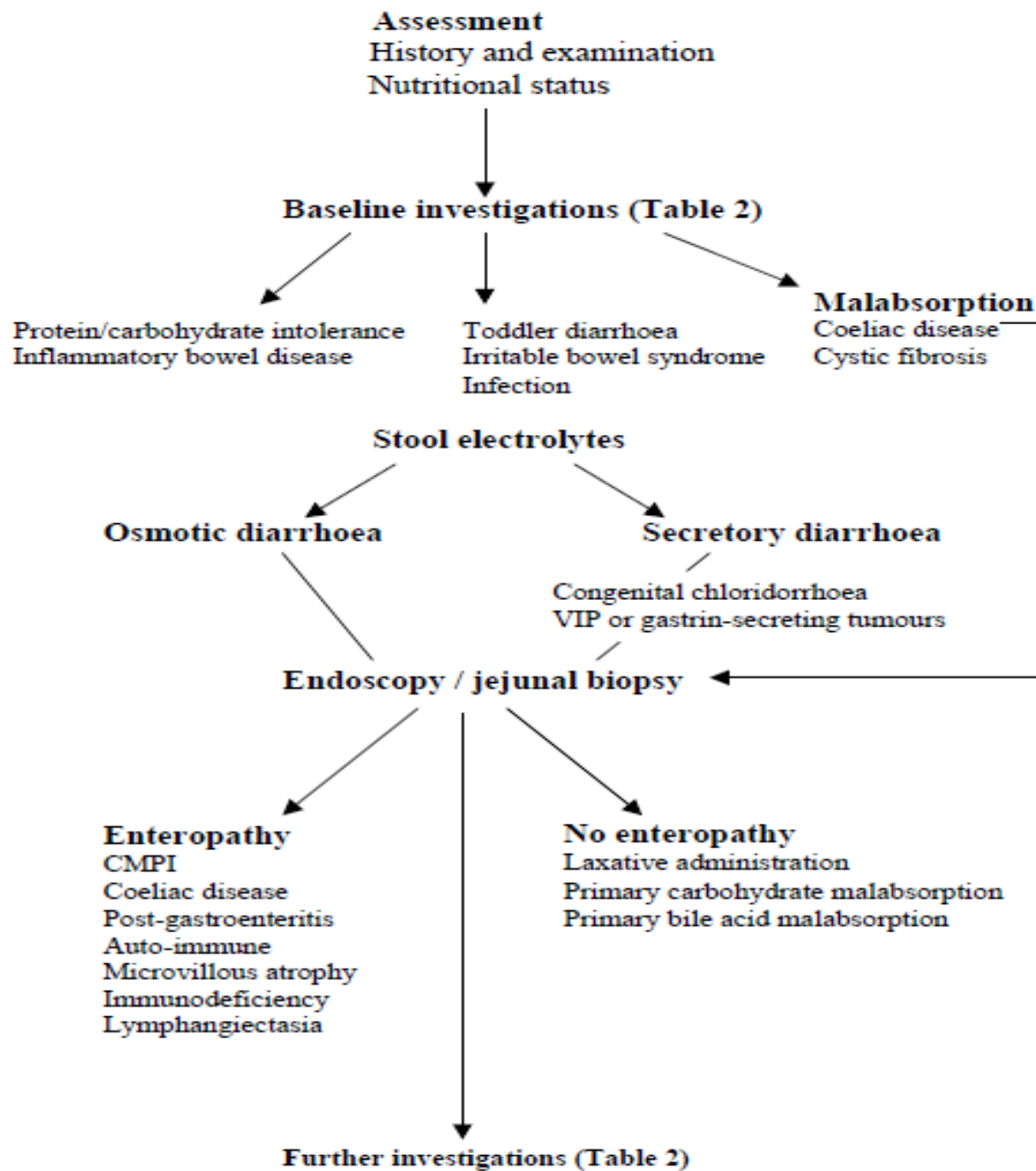
ULCERATIVE COLITIS

- * Onset is adolescence to early adulthood
- * Affects the colon
- * Can have a backwash ileitis
- * Can present with aphthous mouth ulcers
- * Aetiology is poorly understood
- * Treatment steroids, azathioprine and monoclonal antibodies



TB ENTERITIS

- * Can mimic IBD
- * Affects terminal ileum
- * Need tissue to improve diagnostic yield
- * Need to exclude in endemic area
- * Needs 4 drug TB treatment



BASELINE INVESTIGATIONS

Baseline investigations

- Urine microscopy and culture
- Stool: Virology and microbiology
 - Ova cyst and parasites (× 3 fresh samples)
 - Fat globules and chymotrypsin
 - Reducing substances
 - Electrolytes
 - Consider occult blood (× 3, on white diet)
- Full blood count, film and differential, ESR
- Urea and electrolytes, calcium, phosphate, C-reactive protein
- Liver function tests and plasma proteins
- Ferritin

FURTHER INVESTIGATIONS

If possible malabsorption:

- Vitamins B12, A, E and folate
- Clotting screen
- IgA and IgG antigliadin antibodies, anti-endomysial antibodies and/or tissue trans-glutaminase.
- Sweat test
- Hydrogen breath test

Further investigations

- Upper gastrointestinal endoscopy / colonoscopy
- Rectal biopsy
- Abdominal ultrasound
- Barium meal and follow-through
- Technetium-labelled white cell scan
- Abdominal MRI

THANK YOU