Proposed Inpatient department common paediatric cases management

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Pneumonia

Common causative organisms: Streptococcus pneumoniae

Haemophilus influenzae Staphylococcus aureus

Diagnosis:

History:

- Fever
- Cough or difficulty in breathing
- Feeding difficulty

Examination:

- Tachypnoea (increased respiratory rate)
- Signs of respiratory distress (Flaring of the ala nasae, suprasternal, intercostal and subcostal recession)
- Cyanosis
- Weigh the child
- Temperature
- Tracheal position
- Dull on percussion
- Bronchial breath sound, and increased vocal resonance over the consolidation, if any.
- There may have crepitation

Investigations: a) Blood:

Total count of WBC- leucocytosis Differential count of WBC- neutrophilia

- b) X-ray chest: Homogenous opacities or patchy opacities
- c) Blood culture (if possible) may isolate the organism

Treatment:

General:

- Breastfeeding or nutritional support by usual diet
- Keep the baby in well ventilated room
- Clear the blocked nose

Symptomatic:

Paracetamol- 15-20 mg/kg/dose 4-6 hourly if fever is present

Oxygen inhalation if the child has

- -Cyanosis
- -Restlessness
- -Respiratory rate- > 70/min (if age <2mo)

Specific: Antibiotics:

< 2 months of age

Inj. Ampicillin- 200 mg/kg/day in 4 divided dose for 7 days

Inj. Gentamicin- 5 mg/kg single daily dose for 7 days

< 2 months of age

Inj. Ampicilin- 200 mg/kg/day in 4 divided dose for 7 days. If the pneumonia is less severe & the patient can take orally then oral cotrimoxazole or amoxycillin or cepholexin can be given.

or Inj. Procaine penicillin 50,000iu/kg once daily IM for 7 days.

Referral to tertiary care centre / Paediatrician:

Following complications need referral-

- a) Empyemathoracis
- b) Pneumothorax
- c) Unresolved collapse-consolidation
- d) Lung abscess
 - e) Bronchiectasis

Bronchiolitis

Common causative organisms: Respiratory Syncytial Virus (RSV)- Commonest

Parainfluenza virus

Adenovirus

Diagnosis:

History:

- Age 2 months to 2 years, peak age 6 months
- Preceding history of coryza
- Low grade fever
- Feeding difficulty
- Noisy respiration
- Restlessness

Examination:

- Running nose
- Tachypnoea
- Signs of respiratory distress (Flaring of the ala nasae, suprasternal, intercostal and subcostal recession)
- Bloated Chest
- Cyanosis
- Temperature
- Tracheal position
- Hyperresonance on percussion
- Vesicular breath sound with prolonged expiration.
- Wheeze with a few crepitation on the both lung fields
- Liver and spleen may be palpable due to down ward pushing by inflated lungs.

Investigations: a) Blood:

TC of WBC- usually normal

DC of WBC- normal

Hb%- normal

ESR- normal

CRP- (C- Reactive protein)- normal

Oxygen saturation by pulse-oxymetry- low

b) X-ray chest:

Hyper translucency, hyperinflated lung fields

Increased interstitial markings

Horizontal ribs with low flat diaphragm

Treatment:

General:

- Feeding /breastfeeding
- Keep the baby in well ventilated room
- Head up position

Symptomatic:

Paracetamol- 15-20 mg/kg/dose 4-6 hourly if fever is present Nebulization 6 hourly by sulbutamol 0.03ml/kg/dose

+ Ipratropium bromide 250 mg/ml/dose + normal saline 2 ml Clear the blocked nose and OP suction.

Oxygen inhalation

Indication of antibiotic-

- Very sick child
- High fever
- Leucocytosis
- Radiological change, like consolidation

Specific: Antiviral (Ribavirin) in selective cases

Referral to tertiary care centre / Paediatrician:

Following complications need referral-

- a) Very sick child
- b) Need ICU
- c) Combined with collapse-consolidation

Bronchial Asthma

Triggering Factors: Dust

Too hot, too cold foods

House dust mites

Exercise

Animal dander

Pollen

Drugs like aspirin

Diagnosis:

History:

- Recurrent attack of respiratory distress or cough or wheeze (whistling sound arising from chest not from throat or nose)
- Night cough
- Night cough awakening
- Family history of asthma or atopic diseases (allergic rhinitis, allergic conjunctivitis and atopic dermatitis) in the family or atopic diseases of the child.
- Long term cough without any apparent cause

Examination:

- Tachypnoea
- Signs of respiratory distress (Flaring of the ala nasae, suprasternal, intercostal and subcostal recession)
- Cyanosis
- Normal temperature
- Tracheal position
- Hyper resonance on percussion
- Vesicular breath sound with prolonged expiration.
- Bilateral diffuse wheeze

Investigations:

- a) Blood: TC normal, DC may be eosinophilia, Hb%- normal, ESR- normal
- d) X-ray chest:
 - -Hyper inflated lung fields with low flat diaphragm
 - -Tubular heart
- b) Serum IgE- may be high
- c) Lung Function Test (LFT) if age > 8 years

$$\frac{FEV_1}{FVC}$$
 = < 80% of personal best

Reversibility Test – FEV1 > 12 % or 200 ml increased after bronchospasmolysis test

Treatment:

General:

- Breastfeeding and nutritional support
- Head up position

Symptomatic:

Oxygen inhalation

Specific:

Nebulization 6 hourly by sulbutamol 0.03ml/kg/dose of any age and Ipratropium bromide 250mg/ml/dose can be added if the age of the child is more than 2 years

Inj. Hydrortisone 4-5mg/kg/dose 6 hourly for 1-2 days followed by tab. prednisolone 1-2mg /kg/day in 2 divided doses or single morning dose for 3-14 days

Avoidance of triggering factors

Referral to tertiary care centre / Paediatrician:

Following complications need referral-

- a) Persistent bronchial
- b) Refractory asthma
- c) Asthma which needs ICU care

Congenital heart disease

Diagnosis:

History: Recurrent history cough/ difficult breathing

Fatigue on feeding/ exertion

Excessive sweating on feeding and taking longer time to finish feeding

May have bluish colouration of the face or whole body

Not growing well

Examination:

Weigh the child

Clubbing

Tachypnoea

Signs of respiratory distress (Flaring of the ala nasae,

suprasternal, intercostal and subcostal recession)

Conjunctival injection

Bulged praecordium and hyperdynamic

Cyanosis Temperature Tachycardia

Apex beat may be shifted

Thrill may be present

Usually pansystolic murmur can be heart at the praecordium

Basal crepitation

If heart failure, then Tachypnoea

Tachycardia

Cardiomegaly

Enlarged tender liver

Oedema

Investigations:

X-ray chest- Size and shape of the heart

Pulmonary plethora or olegaemia

• ECG- Biventricular hypertrophy

Echocardiogram and colour Doppler- VSD and others.

Treatment

General: Breastfeeding and nutritional support

Symptomatic: Oxygen inhalation

Head up position/ Propped up position

If heart failure then,

Oxygen inhalation Propped up position

Inj. Frusemide 2mg/kg/dose IV Inj. Digoxin calculated dose

Specific: Small VSD close spontaneously other need surgical

intervention

- a) Large VSD, VSD with heart failure
- b) ASD after 2 years
- c) PDA (not responding to medical therapy as in premature baby)
- e) Cyanotic heart diseases eg- TOF

Febrile convulsion

Definition: Generalised convulsion due to rapid rise of temperature resulting from

extra cranial infection.

Diagnosis:

History: Fever

Age: 6 months to 5 years Generalised convulsion

Short duration

Complaints of primary infection (throat pain, drooling

of saliva, frequency of micturition)

Examination:

Weigh the child Temperature: high

Ear, Nose, throat should be examined

Fontanellae: Normal

Signs of meningeal irritation- nuchal rigidity, Kernig sign.

Investigations: a) Blood:

CBC - neutropillic leucocytosis

CRP- raised Urine- R/E & C/S

CXR- may have pneumonia

CSF analysis- should be done in 1st convulsion,

or in doubtful cases.

Blood culture

Treatment:

General: Breastfeeding and nutritional support

Air way clear

Symptomatic: Convulsion is controlled by per rectal Inj. Diazepam

0.5mg/Kg /dose during convulsion and oral Diazepam 0.3- 0.5mg/kg/Day in 3 divided dose for 3-5 days.

Paracetamol-15-20mg/kg/dose orally or per rectally

Tepid sponging

Specific: Antibiotics

Counselling with the parents that it is a benign disease

- a) Complex febrile seizure
- b) Recurrent convulsion
- c) A febrile seizure

Pyogenic meningitis

Common causative organisms: H influenzae

Meningococcus Pneumococus

E coli

Diagnosis:

History: Age of the Child

Fever Convulsion Feeding difficulty

Examination:

Weigh the child Temperature- high Bulged Fontanellae Nuchal rigidity Kernig sign

Investigations

a) Blood:

CBC - neutrophilic lucocytosis

CRP - high.

- b) CXR- consolidation
- c) CSF analysis

Colour- hazy, turbid and rarely clear.

Cell count- nutrophillic leucoytosis (normal 0-5cells/cmm)

Sugar- low (normal 40-80mg/dl) Protein- raised (normal 15-40mg/dl)

Gram stain-

Gram-positive cocci- Pneumococcus

Gram-negative intracellular cocci- Meningococci

Coccobacilli- H influenzae

Bacterial antigen CSF- culture

d) Blood culture

Treatment:

General- Nutritional support

Maintenance of fluid

Symptomatic - Convulsion is controlled by Inj. Diazepam 0.3-0.5 mg/kg/dose per rectum

then maintenance

-Fever: Paracetamol 15-20mg/kg/dose 6 hourly

-Hypoglycaemia: 10% glucose 4 ml/kg/dose I/V then maintenance

Specific:

Inj. Ampicillin 400mg/kg/day in 4 devided doses plus Inj. Chloramphenicol 100mg/kg/day in 4 divided doses or

Inj. Ceftiaxone 100mg/kg/dose single daily dose. Antibiotic should be continued until 5 days after abatement of fever.

Inj. Dexamethasone 0.15 mg/kg/dose 6 hourly for 2 days

- a) Hydrocephalus: drug treatment and Surgical consultation
- b) Palsy: Physiotherapy
- c) Epilepsy: Antiepileptic drugs
- d) Subdural effusion: May need subdural tap
- e) Mental retardation
- f) Learning deficit
- g) Vision and hearing problem

Encephalitis/Cerebral malaria

Common causative Organisms:

Virus- Herpes Virus, mumps, measles virus

Parasites- Malarial parasites

Diagnosis:

History:

Unconsciousness of recent onset Focal or generalised convulsion High fever in case of malaria in the malarial area

History of coryza, malaise, myalgia, bodyache History of poisoning, drug ingestion or trauma.

Examination:

Varying grade of unconsciousness: drowsy/ stupor/ coma

Decorticated/ cerebrated posture

Herpetic vesicles or lesion usually in the face

Usually high temperature Pupil size and reaction to light

Cranial nerve involvement

Limbs- sensory/ motor involvement

Signs of meningeal irritation- Nuchal rigidity, kernig sign

Investigations: a) Blood

Blood count- normal

Blood film-thick & thin film to identify malaria

- b) Urine R/M/E & C/S
- c) X-ray chest
- d) CSF analysis:

Colour- clear.

Cell count- normal (normal 0-5cells/cmm)

Sugar- normal (normal 40-80mg/dl) Protein- normal (normal 15-40mg/dl)

Gram stain- no bacteria present Bacterial antigen- negative

CSF- culture- negative

- g) Electrolytes- delusional hypornatraemia
- h) Blood Sugar, Blood urea, S. creatinine

Treatment:

General:

- Maintain nutrition by IV fluid or NG feeding
- Changes of posture 2 hourly
- Care of bowel and bladder

Symptomatic:

Paracetamol to reduce fever Diazepam to control convulsion IV glucose to prevent hypoglycaemia

Specific:

Inj. Jasoquine 20mg/kg bolus IV in isotonic fluid (5-10ml/kg) over 3-4 hour then 10mg/ Kg /dose IV 8 hourly until consciousness regain and then oral Jasoquine 10mg / Kg/dose 8 hourly, total 5-7 days

Inj. Acyclovir 10 mg /Kg/ dose IV 8 hourly for 10 days

- a) Focal paresis/ palsy- Physiotherapy
- b) Epilepsy: Antiepileptic drugs
- c) Mental retardation
- d) Learning deficit
- e) Vision and hearing problem
- f) Behavioural problem

Enteric Fever

Causative organism: Salmonella typhi and Salmonella para typhi

Diagnosis:

History: Fever

Headache Poor feeding Abdominal pain

Examination:

Weigh the child
Toxic look
Coated tongue
Truncal rash
High temperature
Abdominal tenderness
Hepatosplenomegaly

Liver dullness may be obliterated

Bowel sound

Investigations: a) Blood:

TC of WBC- leucopenia

DC of WBC- relative lymphocytosis

Blood Culture- in 1st week Widal test - beyond 1st week Urine for R/M/E & C/S CXR- consolidation.

Treatment:

General:

Nutrition support

• Care of mouth

Symptomatic: Paracetamol to reduce temperature

IV fluid

Specific:

Antibiotic:

Cotrimoxazole- 5 mg/kg/dose in term of trimethoprim 12 hourly

for 14 days.

Ciprofloxacin 10mg/kg/dose 12 hourly for 14 days Ceftriaxone 100mg/kg/single daily dose for 10 days

Cefixim 10mg/kg/dose 12 hourly for 10 days

- a) Perforation-Nothing per oral
 - -IV fluid antibiotic
 - -Surgical consultation
- b) Carrier state if require surgery
- c) Other complications

Prevention:

Hand washing before taking food and after coming from toilet.

Safe water drinking

Vaccination

Treatment of carrier state:

Amoxicillin 50mg/kg/day in 3 divided dose for 10 days May require cholecystectomy

Urinary Tract Infection (UTI)

Common causative organism: E coli

Diagnosis:

History: Fever

Frequency of micturition Painful micturition Abdominal pain Not growing well

Examination:

Weight Height

Temperature may be raised Supra pubic tenderness Tender renal angle

Investigation:

Blood count

Urine R/M/E and C/S with colony count

Treatment:

General:

- Nutrition support
- Plenty water to drink

Symptomatic: Paracetamol to reduce temperature

Regular evacuation of bowel habit

Specific:

Antibiotic:

Cotrimoxazole- 5 mg/kg/dose in term of trimethoprim 12 hourly for 7 days.

Cephradine 50mg/ Kg/day in 3 divided dose for 7days Ciprofloxacin 10mg/kg/day 2 divided dose for 7days

- a) Recurrent UTI
- b) UTI below the age of 5 years

Severe Malnutrition

Diagnosis:

History: Detailed dietary history

Milk formula feeding

Early weaning/late weaning Diluted formula feeding

Discontinuation of breast-feeding

Recent history of diarrhoea, pneumonia and other infections

(measles, pertussis, Tb)

Poor illiterate parents

Examination: Take weight and height/length of the child

Growth retardation that is low weight for age or

weight for height.

Wasting

With or without oedema

Pallor

Glossitis, angular stomatitis

Hair change- Sparse, lustreless, thin, easily pluckable

hair, flag sign.

Eye changes- Conjunctival xerosis, Bitot spot, corneal xerosis,

corneal, ulceration and opacities.

Skin changes- cold clamy skin, loose nonelastic, hanged skin specially

at buttock. hyperpigmentation, hypopigmentation

desquamation and ulceration

Hypothermic, sometimes features of shock.

Investigations:

a) Blood

CBC – neutrophillic leucocytosis if infection

Hb- will be low

ESR-high

- b) Urine for R/M/E and C/S
- c) Blood sugar level- low (<3mmol/L in case of hypoglycaemia)
- d) X-ray chest: microcardia
- e) MT

Treatment:

- 1) Immediate resuscitation to combat-
 - -Hypothermia: by covering the body

keep the child in warm wrapping with cloths/by using fan heater

-Hypoglycaemia: 4ml/Kg of 10% glucose IV stat or 50 ml 10% glucose

through nasogastric tube where IV administration not possible.

Early feeding

- -Dyselectrolytemia: ORS by mouth or ReSoMal by mouth
- -Vitamin A deficiency: If age < 6month then 50,000 iu once

6-12 month then 1 lac unit once

>12 month then 2 lacs unit once

One dose for malnutrition and three dose if eye sign of vitaminA deficiency present.

-Control of infection by antibiotic, inj. ampicillin 200mg/kg/day + inj. gentamicin 5mg/kg/day

-Severe anaemia (<4mg% of Hb) needs blood transfusion

2) Feeding-

Round the clock calculated nasogastric tube feeding should be started as early as possible.

Fluid- 100ml/kg/day initially and can be increased later on

Calorie- 110kcal/kg/day

2 hourly 12 divided feeding

Preparation: For each feeding milk should be given 1 teaspoonful to 1 ounce (30 ml) water, rest of the calorie should be calculated by adding sugar and soyabin oil (1 tsf powder milk=whole cows milk= 20 kcal, 1 tsf sugar= 20 kcal and 1 tsf soyabin oil= 45 kcal)

Nasogastric tube should be withdrawn if the child can take three-quarter of calculated feed by orally.

3) Micronutrients supplementation

- Potassium: 3mEq/Kg/day (1 tsf= 10mEq)
- Zinc: 1-2mg/Kg/day (1 tsf= 10mg)
- Magnesium: 2-3 mg/Kg/day (1 tsf= 50mg)
- Multivitamin can be given daily 1 ml twice daily
- Iron should be given (6mg/Kg/day) from 2nd week of initial phase.
- 4) Mother or care giver should be taught regarding feeding to be continued at home
- 5) If not possible to follow above management then refer the child to tertiary care centre

Haemolytic Anaemia

Diagnosis:

History: Age- usually above 6 months

Consanguinity between parents

Progressive pallor

Previous history of blood transfusion

Not growing well.

Examination: weight

Height Pallor Jaundice

Thalassemic facies (frontal bossing, depressed nasal bridge,

malar prominence and malocclusion of teeth)

Hepatosplenomegaly

Investigation:

Blood count- Normal

Hb%-Low

Blood film- Anisopoikilocytosis with

hypochromic-microcytic RBC's. There are some schistocyte, target cells, nucleated RBC's and a few macrocytes. WBC's are mature and normal count.

Platelets count normal.

X-ray of hand and skull lateral view

Serum Iron

Serum ferritin, TIBC Osmotic fragility test

Haemoglobin electrophoresis

Treatment:

General:

Nutrition support Avoid iron rich food Zinc supplementation

Symptomatic: Repeated packed blood cell transfusion

Iron chelating agent Inj. Desferol

Antibiotic if infection

Specific:

Bone-marrow transplantation

Referral to tertiary care centre/Paediatrician:

a) To establish diagnosis

b) To assess endocrine function

Iodine Deficiency Disorder (IDD)

Diagnosis:

History: Geographical location, mountainous areas, family history swelling of the anterior aspect of the neck

Examination: Palpation of goitre

Table: Classification of goitre (WHO classification)

Grade 0	No palpable or visible goitre
Grade 1	A mass in the neck that is consistent with an enlarged thyroid that is palpable but not visible when the neck is an normal position. It moves upwards in the neck as the subject swallows
Grade 2	A swelling in the neck that is visible when the neck is in normal position and is consistent with an enlarged thyroid when the neck is palpated

Investigations:

- USG of the thyroid gland
- Serum T₃, T₄, TSH estimation
- Radioiodine uptake
- Urinary iodine excretion (normally 90% iodine is excreted through urine)

Treatment

Iodized salt

Sodium-L-thyroxine

Nephrotic Syndrome

Diagnosis:

History: Age 2 - 10 years

Generalised swelling

Scanty normal coloured urine

May have recurrent history of swelling

Examination:

Weight

Body surface area (Wt in Kg \times 4 + 7/90+ Wt in Kg)

Blood pressure- normal

Oedema, including scrotal/labial swelling

Ascites Anasarca

Investigations:

a) Urine- Appearance/colour- normal or frothy

Albumin- +++ (massive proteinurea)

RBC- nil

Pus cell- may be present

Cast- hyaline cast may be present

Urine- culture and coloney count

UTP- >1gm/m²/day (Nephrotic range of proteinurea)

b) Protein: Creatinine ratio- > 3 indicate nephrotic range

of proteinurea

- c) Serum Albumin= < 2.5 gm/dl
- d) S. Cholesterol = > 220 gm/dl
- e) S. Electrolytes
- f) Blood count, blood urea, creatinine/ BUN
- g) CXR- to exclude pulmonary infection/ effusion
- h) HB_SAg
- i) CBC and ESR

Treatment:

General:

- Diet: liberal diet/salt restricted
- Maintain intake-output chart, albumin and weight chart

Symptomatic:

For huge ascitis: diuretics – spironolactone and frusemide 25% human albumin 0.8- 1g/kg IV

• If infection (UTI, peritonitis, septicaemia, skin infection)

- antibiotics

• Abdominal pain with fever

Inj. Penicillin 2 lac/Kg/day in4 divided dose for 7 days

• H₂ blocker/ antacid during steroid therapy

Specific:

- Standard Prednisolone therapy (SPT): 60mg/m²/day in 3 devided dose for 6 weeks followed by 40mg/m² alternate day single morning dose for 6 weeks.
- Counselling:
 - -Recurrent attack
 - -No deterioration of kidney function
 - -Good prognosis. There is high chance of cure at the end of second decade

- a) Nephrotic syndrome with hypertension, haematuria and infection
- b) If age below 2 years and above 10 years
- c) No response to standard prednisolone therapy

Acute glomerulonephitis (AGN)

Mainly post streptococcal (either skin infection or throat infection) sequelae.

Diagnosis:

History: Scantly high colour urine

Puffiness of the face

Swelling of the whole body May have- Headache

Vomiting

Blurring of vision Breathing difficulty

Convulsion

Examination:

Weight

Body surface area (Wt in Kg \times 4 + 7/90+ Wt in Kg)

Blood pressure- Usually high

Puffiness of the face or eyelid swelling

Oedema

May have heart failure

Tachycardia Tachypnoea

Enlarged tender liver Basal crepitation

Impairment of consciousness.

Investigations:

a) Urine for R/M/E

Smoky or coco cola colour Albumin present- +/++ Plenty RBC and RBC cast

A few pus cells.

- b) Blood urea/BUN- raised
- c) Serum creatinine- raised
- d) X-ray chest- may have cardiomegaly pulmonary oedema.
- e) ECG- hyperkalaemia
- e) ASO titre may be raised

Treatment

General: -Bed rest

-Salt and protein restricted

-Potassium and potassium containing foods and fruits restricted.

-Restriction of fluid 400 ml/m²/day + previous day output

-Maintain intake-output chart, albumin, weight and BP chart

Symptomatic: Hypertension – Diuretics and/or antihypertensive drugs

Heart failure - a) Diuretics

b) Digoxin

c) Bed rest

Specific: -Tab. penicillin 50mg/Kg/ day in 4 divided dose for 10 days.

- a) Acute renal failure
- b) Heart failure
- c) Volume overload
- d) Hyperkalaemia
- e) Hypertensive encephalopathy