





Department of Health Research Ministry of Health and Family Welfare, Government of India



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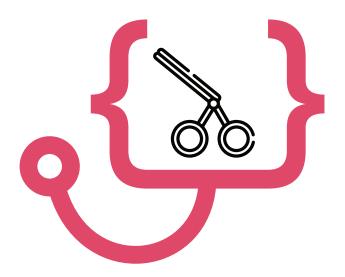
world Health Organization

Suggested Citation: Standard Treatment Workflows of India, 2019 Edition, Vol. 1, New Delhi, Indian Council of Medical Research, Department of Health Research, Ministry of Health and Family Welfare, Government of India

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Printed in India









Department of Health Research Ministry of Health and Family Welfare, Government of India



सबका साथ, सबका विकास, सबका विश्वास Sabka Saath, Sabka Vikas, Sabka Vishwas



डॉ हर्ष वर्धन **Dr Harsh Vardhan**

स्वास्थ्य एवं परिवार कल्याण, विज्ञान और प्रौद्योगिकी व पृथ्वी विज्ञान मंत्री, भारत सरकार

Union Minister for Health & Family Welfare Science & Technology and Earth Sciences Government of India



FOREWORD

As per National Health Policy 2017, Health for all which is equitable, accessible and affordable is a priority for Government of India. The health sector in the country is working hard towards achieving Universal Health Coverage and Sustainable Development Goals. As the transition to value-based care moves forward, the focus in patient care is shifting to both quality and quantity.

2. Providing quality healthcare is a key focus area. I congratulate Department of Health Research and Indian Council of Medical Research for addressing this issue through the development of Standard Treatment Workflow that are evidence based as well as locally contextual with the aim of guiding decisions and criteria regarding diagnosis, management, and treatment of common diseases. These workflows will play a pivotal role in realizing the dream of Universal Health Coverage with equitable health systems.

I am happy to state that within a year of initiating this ambitious activity, the first 3. volume is ready to be released. I am confident that all stakeholders contributing towards healthcare in the Country will benefit from these workflows.

Congratulations to the Team!

Jaroh

(Dr. Harsh Vardhan)

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प्रोफेसर (डा.) बलराम भार्गव, पदम श्री

एमडी, डीएम, एफआरसीपी (जी.), एफआरसीपी (ई.), एफएसीसी, एफएएचए, एफएएमएस, एफएएससी, एफ.एन.ए., डी.एस.सी.

सचिव, भारत सरकार स्वास्थ्य अनुसंधान विभाग स्वास्थ्य एवं परिवार कल्याण मंत्रालय एवं महानिदेशक, आई सी एम आर

Prof. (Dr.) Balram Bhargava, Padma Shri

MD, DM, FRCP (Glasg.), FRCP (Edin.), FACC, FAHA, FAMS, FNASc, FASc, FNA, DSc

Secretary to the Government of India Department of Health Research Ministry of Health & Family Welfare & Director-General, ICMR



भारतीय आयुर्विज्ञान अनुसंधान परिषद स्वास्थ्य अनुसंधान विभाग

स्वास्थ्य एवं परिवार कल्याण मंत्रालय भारत सरकार वी. रामलिंगस्वामी भवन, अंसारी नगर नई दिल्ली - 110 029

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Preface

To revive clinical medicine and encourage rational use of drugs, diagnostics and other healthcare services, we embarked upon this ambitious initiative of developing standard treatment workflows for common and serious diseases encountered by the treating doctors at all levels of health system.

In addition, we also wanted to create a robust pre-defined referral mechanism to decongest the tertiary centres and revitalize the primary and secondary health facilities.

I hope these resource stratified treatment workflows would be useful to doctors working at primary, secondary as well as tertiary levels and help in optimal utilization of resources not only in our country but also other countries like ours where the resources are limited and demands on the health system are high.

Balsone She

(Balram Bhargava)

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SPECIALITIES COVERED IN THIS EDITION

- CARDIOLOGY

ATRIAL FIBRILIATION BRADYARRTHYMIAS HEART FAILURE STABLE ANGINA STEMI LINSTARI E ANGINA/ NS

- EN1

ACUTE RHINOSINUSITIS CHRONIC RHINOSINUSITIS EPISTAXIS HEARING IMPAIRMENT IN PEDIATRIC AGE GRC NECK SPACE INFECTION OTORRHOEA PHARYNGITIS AND SORE THROAT

- NEPHROLOGY

ACUTE KIDNEY INJURY CHRONIC KIDNEY DISEASE GLOMERULONEPHRITIS URINARY TRACT INFECTION

- NEUROLOGY

APPROACH TO ACUTE PARALYSIS DEMENTIA EPILEPSY HEADACHE NEUROINFECTIONS STROKE

- OBG

ANTENATAL MANAGEMENT DILATATION AND CURETTAGE HEAVY MENSTRUAL BLEEDING HYSTERECTOMY POSTPARTUM HAEMORRHAGE UTERINE FIBROIDS AND POLYPS

- PAEDIATRICS

ACUTE DIARRHEA DENGUE FEVER FEVER IN CHILDREN SEPSIS AND SEPTIC SHOCK IN CHILDREN SEVERE ACUTE MALNUTRITION SEVERE PNEUMONIA IN CHILDREN

- PSYCHIATRY

ALCOHOL USE DISORDERS ANXIETY DISORDERS CHILDHOOD BEHAVIOURAL DISORDERS CHILDHOOD EMOTIONAL DISORDERS CHILDREN WITH DEVELOPMENTAL DISORDERS DEPRESSION PSYCHOSIS SOMATOFORM DISORDERS

- PULMONOLOGY

ACUTE RESPIRATORY INFECTION ASTHMA CHRONIC OBSTRUCTIVE PULMONORY DISORDER RESPIRATORY FAILURE

- UROLOGY

ACUTE URINARY RETENTION IN ME GROSS HAEMATURIA MALE INFERTILITY RENAL AND URETRIC STONES SCROTAL SWELLING

• CONTRIBUTORS



INTRODUCTION

GOAL

To empower the primary, secondary and tertiary care physicians/surgeons towards achieving the overall goal of Universal Health Coverage with disease management protocols and pre-defined referral mechanisms by decoding complex guidelines

OBJECTIVES

Primary Objective:

To formulate clinical decision making protocols for common and serious medical/ surgical conditions for both OPD and IPD management at primary, secondary and tertiary levels of healthcare system for equitable access and delivery of health services which are locally contextual

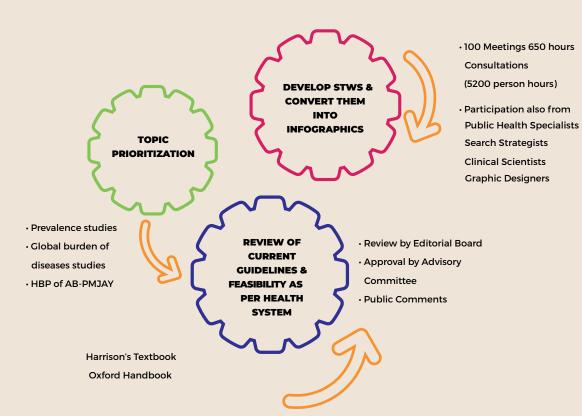
Secondary Objective:

To facilitate PMJAY arm of Ayushman Bharat with secondary and tertiary level management of all surgical and medical conditions covered under the scheme.

METHODOLOGY











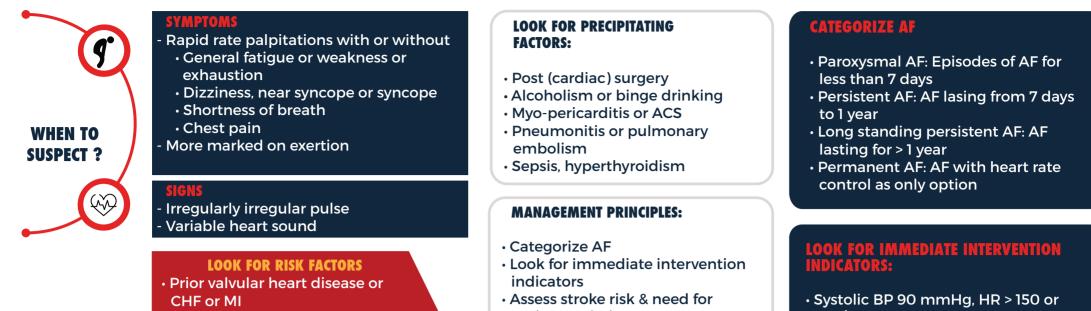






Standard Treatment Workflow (STW) for the Management of **ATRIAL FIBRILLATION**

ICD-10-148.91



- Prior TIA or stroke or embolic episode
- Hypertension, DM, COPD, CKD, Obesity
- anti-coagulation
- Assess bleeding risk
- Need for rate control
- Consideration for rhythm control
- <50/min
- Ongoing Angina
- CHF or TIA or stroke
- Major bleed on Oral Anti-coagulants

STROKE RISK SCORE

BLEEDING RISK SCORE

CHA ₂ DS ₂ -VAS _C	SCORE	HAS-BLED	SCORE
- <u>C</u> ongestive heart failure/LV dysfunction	1	- Hypertension i.e. uncontrolled BP	1
- <u>Hypertension</u>	1	- Abnormal renal/liver function	1 or 2
- <u>Aged ></u> 75 years	2	- Stroke	1
- <u>D</u> iabetes mellitus	1	 Bleeding tendency or 	1
- <u>S</u> troke/ TIA/ TE	2	predisposition	1
- <u>V</u> ascular disease [prior MI, PAD or aortic	1	- Labile INR	1
plaque]		- Age (e.g. >65)	
- <u>Ag</u> ed 65-74 years	1		1
		- Drugs (e.g. concomitant aspirin or	
- <u>S</u> ex category [i.e. female gender]	1	NSAIDSs or alcohol	
Maximum Score	9		9

Bleeding Risk High in score >3

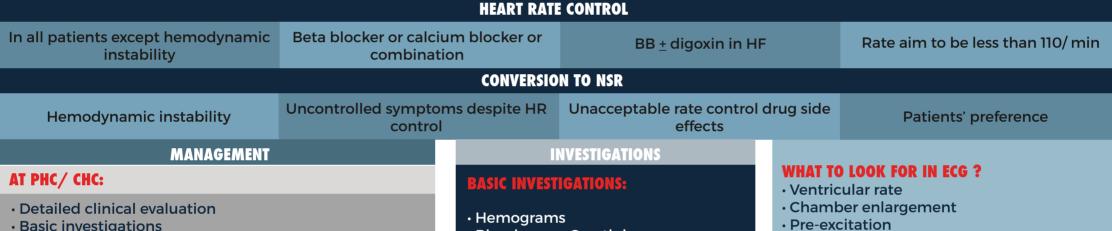
CHOICE OF ANTI-COAGULATION:

Vitamin K antagonist

- Aim for INR 2-3
- Assess risk of bleeding
- Take measures to reduce/ modify risk of bleeding
- Dietary modification & regular monitoring

MEASURES TO REDUCE HIGH BLEEDING RISK:

- Control SBP to less than 140 mmHg
- Avoid dietary indiscretions
- Avoid concomitant aspirin, anti platelets, NSAIDs
- Avoid alcohol
- Correct anemia



Basic investigations

- - oatini

- Careful ECG evaluation
- Start OAC if indicated (based on Stroke risk)
- Start Metoprolol if HR >110/ min & no evidence of CHF
- Refer if indicators for early intervention

OAC if score >1 in men and >2 in women

AT DISTRICT HOSPITAL:

- Admit if indicators of early interventions
- Immediate cardioversion after heparinization, if hemodynamic instability
- Manage precipitating factors if any
- · Assess stroke, bleeding risk & coagulation parameters
- Detailed echocardiogram
- Start OAC, maintain INR around 2-3
- Control HR by single drug or combination of BB & Ca **Blocker**

Refer HR uncontrolled or CHF or angina

AT TERTIARY CENTRE:

Yes

Re-assess clinical status, adequacy of AC

Yes

Drug version

Successful

Yes

Continue drug

No

DC version

- Consider need of NOAC
- Optimise management of underlying cardiac disease
- Stress life style and AF risk factor modification
- Assess need for rhythm control and discuss pros & cons
- Consider RFA in select patient

- Electrolytes
- 12 lead ECG

DESIRABLE INVESTIGATIONS:

- Plain X-ray chest
- Thyroid evaluation
- Liver function test
- Troponins
- Prothrombin time, INR (Coagulation profile)
- Echocardiography

OPTIONAL INVESTIGATIONS:

- Prolonged ECG monitoring
- Trans-esophagial echocardiography

No

No Very rapid HR >130/min

Yes

No Symptomatic

No

Clinical

Follow-up

- Exercise Stress Test
- CT scan
- MRI

Б Yes

CHF

HR control

Aim <110/min

HR > 110/ min

Consider DC version

No

BB or Ca Block

or combine

Yes

Careful BB/Dig

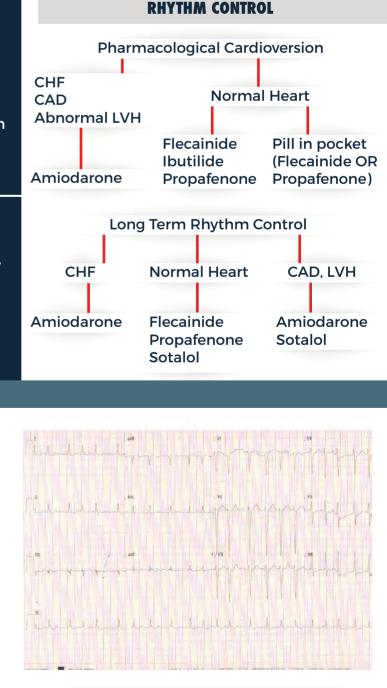
Amiodarone

Sign/ symptoms suugestive of AF Confirm by 12 channel rhythm strip Hemodynamic Instability

- EP study
- Coronary angiography



- Bundle branch block
- QT interval



Anti-coagulants in all Except

• Reversible

Score <1 (men); <2 (women)

🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

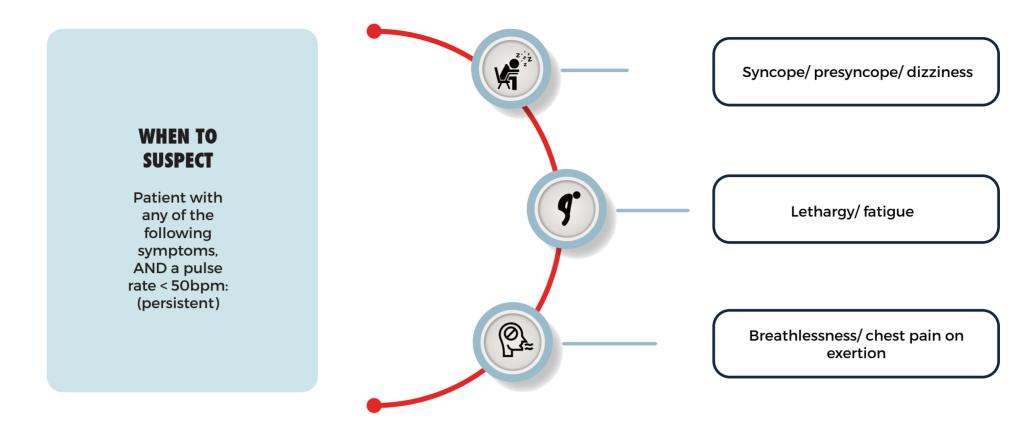
This STW has been prepared by national experts of India with feasibility considerations for various levels of healthcare system in the country. These broad guidelines are advisory, and are based on expert opinions and available scientific evidence. There may be variations in the management of an individual patient based on his/her specific condition, as decided by the treating physician. There will be no indemnity for direct or indirect consequences. Kindly visit our web portal (stw.icmr.org.in) for more information. © Indian Council of Medical Research and Department of Health Research, Ministry of Health & Family Welfare, Government of India.

MANAGEMENT ALGORITHM





Standard Treatment Workflow (STW) for the Management of BRADYARRTHYMIAS IN SYMPTOMATIC PATIENTS ICD-10-R00.1



BASIC EVALUATION

HISTORY

- Syncope/ presyncope: frequency, associated fall/injury/incontinence
- Exertional angina or known coronary artery disease
- Known hypothyroidism or kidney disease
- On beta-blockers, Calcium Channel Blockers or digoxin
- $\boldsymbol{\cdot}$ Patient with an implanted pacemaker or other device
- Yellow oleander poisoning

EVALUATION AND TREATMENT OF UNSTABLE PATIENTS

1. TREATMENT OF ASSOCIATED CONDITIONS

- Hyperkalemia
- Suspected drug (BB or CCB) overdose:
- i. Withhold the drug
- ii. iv insulin (1 U/kg bolus followed by 0.5 U/kg/h) with glucose monitoring(or) iv glucagon if available
- 2. TEMPORARY PACEMAKER INSERTION

(iv dopamine or adrenaline may be given till the time TPI can be placed)

EXAMINATION

- Drowsiness/ impaired consciousness
 BP, heart rate
- **TESTS TO BE DONE**

Patient presenting to PHC/CHC:

- 12-lead ECG
- Blood urea, serum creatinine
- Electrolytes
- Blood sugar

EVALUATION AND MANAGEMENT OF STABLE PATIENTS

Findings on 12-lead ECG

- Atrioventricular block
- Sinus node dysfunction
- Other conduction disorders with 1:1 AV conduction
- Non-diagnostic ECG

INDICATIONS FOR URGENT TREATMENT/REFERRAL

GENERAL APPROACH TO PATIENTS WITH SYMPTOMATIC BRADYCARDIA

• Hypotension (SBP <90 mmHg), impaired consciousness or

1. Rule out **associated conditions**

October/ 2019

N

- ongoing chest pain
- Recurrent or ongoing syncope/presyncope
- Associated headache with or without neurologic deficit (suspect intracranial event)
- Patient with a pre-existing device
- If ECG available, evidence of any of the following
 - Complete heart block
 - Sinus node disease with pauses >3 s long
 - Bradycardia (HR < 50 bpm)
 - (with or without hyperkalemia, serum K > 5 mEq/L)

RECOMMENDED PACING MODES

- Atrial-based single or dual chamber pacing

- VVI pacing is reasonable if symptoms are

- VVI/Dual chamber pacing in patients with

- CRT (or HBP) in patients with LVEF 36-50% and requiring ventricular pacing

- Renal dysfunction, hyperkalemia
- Drug toxicity (BB, CCB, clonidine, Lithium)
- Sleep apnea (clinical scoring systems such as Epworth Sleepiness Scale may be used for initial assessment)
- 2. Transthoracic echocardiography

INDICATIONS FOR PERMANENT PACING

AV NODAL DISEASE

 Complete heart block, advanced AV block, or Mobitz Type II block

1. SND with intact AV conduction

infrequent

LVEF >50%

>40% of the time

- CRT (or HBP) if LVD <35%

2. AV node disease

- Symptomatic patients with AV block other than above
- Associated neuromuscular disease

Symptomatic patients with sinus pauses > 3 s long with symptom correlation

SINUS NODE DYSFUNCTION

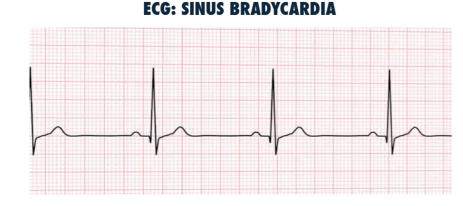
 Asymptomatic patients with sinus pauses > 6 s long

OTHER CONDUCTION DISORDERS WITH 1:1 AV CONDUCTION

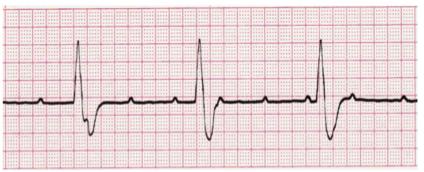
Symptomatic patients with HV ≥100 ms on EPS
Others (alternating BBB, infiltrative/ neuromuscular disease)

ADDITIONAL TESTING

- 1. **Advanced imaging** (cMRI) may be needed if infiltrative disease is suspected
- 2. Ambulatory ECG may be needed
 - In patients with first or second degree AV block for symptom correlation
 - In patients with suspected sinus node disease for detection of pauses and symptom correlation
 - In symptomatic patients with LBBB or bifascicular block
- 3. Implantable Loop Recorder and EPS (consult published society guidelines)



ECG: THIRD DEGREE HEART BLOCK



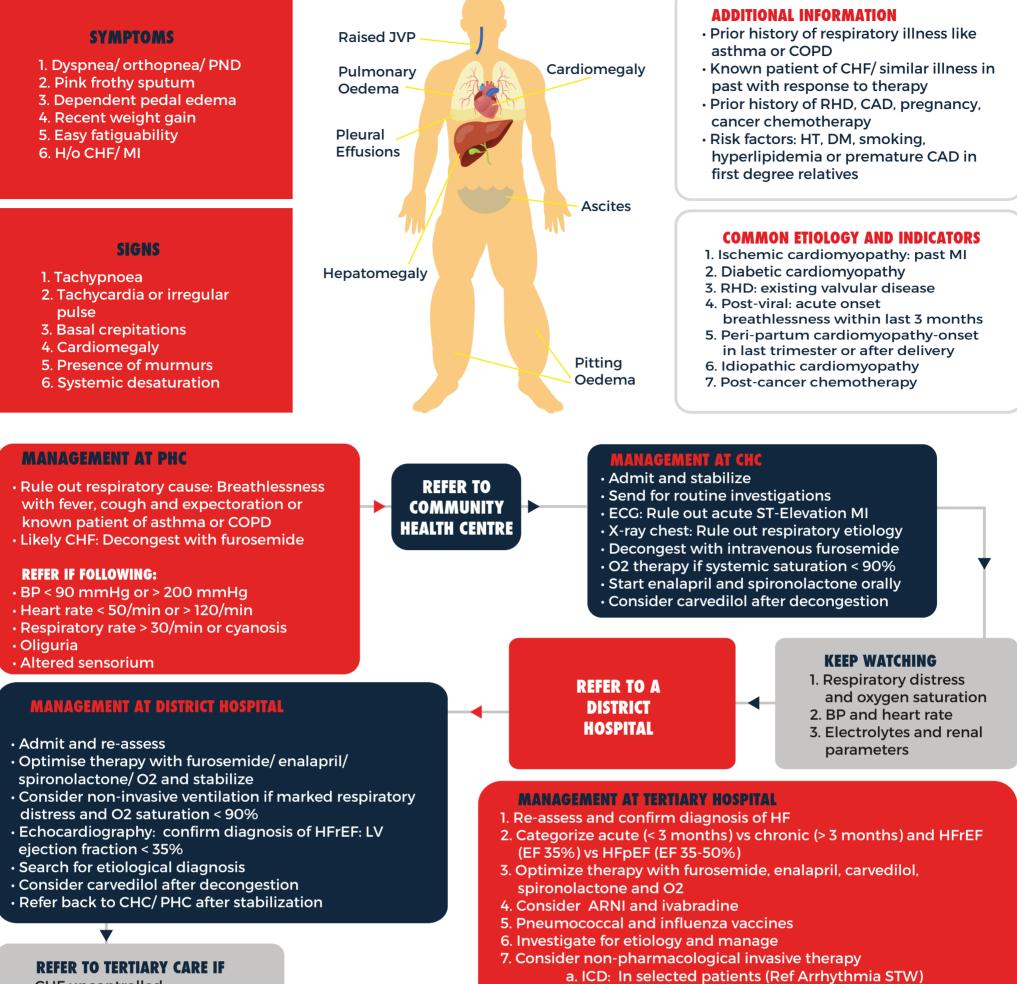






Standard Treatment Workflow (STW) for the Management of **HEART FAILURE: A BREATHLESS PATIENT**

ICD-10-150.9



CHF uncontrol Unstable hemo Suspected ong Abnormal elec Abnormal rena Structural hear Unclear etiolog	odynamics going ischemia trolytes al functions rt disease			b. BiV: EF <35 morph 8. Etiology based a. PCI	Consider in N 5%, QRS >1506 nology and o d Interventio replacement	msec in sinus rh ptimal medical ns	Symptomatic patient , ythm with LBBB therapy of >3 months
			CONSIDE	R AT ALL LEVELS			
Smoking Cessation		ysical tivity	Weight Reduction	Moderation of alcohol	Control of D HTN/ Lipic	PM/ with	ndary CVD prevention n aspirin and statins
			INVES	TIGATIONS:			
 BASIC INVESTIGATIONS Hemogram, ESR Blood sugar Urine examination Urea/ Creatinine Sodium/ Potassium ECG Chest X-ray PA view 	Hemogram, ESR Blood sugar Jrine examination Jrea/ Creatinine Sodium/ Potassium ECG · · · Cardiomegaly · Pulmonary venous congestion · Pneumonia or other		 WHAT TO LOOK FOR IN AN ECG? Pathological Q wave Conduction abnormalities, especially LBBB Chamber enlargement Atrial fibrillation Note: If ST elevation present, manage as STEMI 		DESIRABLE INVESTIGATIONS • 2D Echocardiography • BNP/NT pro-BNP • Troponin • Lipid profile • Thyroid function test • Iron profile		 OPTIONAL INVESTIGATION Prolonged ECG monitoring Coronary angiography Radionuclide imaging CT scan MRI PET Myocardial biopsy Electrophysiological study
			COMMON DRUG	GS AND DOSAGE FOR	CHF		
FUROSEMIDE • Dose 20-80 mg daily PO • Intravenous 10-40 mg SO • Change to oral when syr • Monitor serum electroly on therapy	OS in acute stage mptoms subside	ric acid	 Start after d with BP > 10 Uptitrate do tolerable do 	o 25 mg twice daily lecongestion with le 00 mmHg and HR > ose 1-2 weekly till m ose 1 on BP, heart rate a	ow dose 60/ min aximum	• Start with mmHg, no creatinine	0 10 mg twice daily PO low dose with BP >100 ormal electrolyte and less than 2.5 mg/dl lose 1-2 weekly till maximum
 SPIRONOLACTONE Dose 25-50 mg once dai Keep watch on serum pevery 2-4 weekly 	-	iine	recipitation • Increase div	of CHF symptoms aretics and reduce of reappearance of CH	carvedilol	 Keep watc 	h on BP and electrolytes ry increment and on
		🖝 KEEP	A HIGH THRESH	OLD FOR INVASIVE	PROCEDURE	s	
ABBREVIATIONS	overter defibrillator	CABG:	ercutaneous Corc Coronary Artery Cardiovascular Di	•••	HFpEF: H	leart Failure wit	h reduced Ejection Fraction h preserved Ejection Fraction

BiV: Bi-Ventricular Pacing **PND:** Paroxysmal Nocturnal Dyspnea **RHD:** Rheumatic Heart Disease **CAD:** Coronary Artery Disease

STEMI: ST elevation Myocardial Infarction **LV:** Left Ventricle **COPD:** Chronic Obstructive Pulmonary Disease

REFERENCES

1. Management Protocols for Chronic Heart Failure in India. Mishra S, Mohan JC, Nair T et al. Indian Heart J.2018;70:105-127,

2. ESC Guidelines for the diagnosis and treatment of acute and chronic heart failure: The Task Force for the diagnosis and treatment of acute and chronic heart failure of the European Society of Cardiology (ESC). Developed with the special contribution of the Heart Failure Association (HFA) of the ESC. Ponikowski P, Voors AA. Anker SD et al. European Heart Journal. 2016;37:2129-2200

- 3. Chronic heart failure in adults: diagnosis and management. NICE guideline [NG106] Published date: September 2018
- 4. 2013 ACCF/AHA Guideline for the Management of Heart Failure. A Report of the American College of Cardiology Foundation/American Heart Association Task Force on Practice Guidelines. Yancy CW, Jessup M, Bozkurt B. J Am Coll Cardiol. 2013;62-16: e150-e210





Standard Treatment Workflow (STW) for the Management of **STABLE ANGINA**

ICD-10-120.9

PATIENT PRESENTING WITH CHEST PAIN



CONSIDER ANGINA IF

- Diffuse retrosternal pain, heaviness or constriction, radiating to arms or neck or back
- Associated with sweating
- Easily reproduced with post-meal exertion
- Consider atypical presentation: Exertional fatigue or breathlessness or profuse sweating or epigastric discomfort

Likelihood more if known patient of CAD

ANGINA UNLIKELY IF

- Variable location or characteristic
- Long lasting (hours to days) or short lasting (less than a minute)
- Restricted to areas above jaw
 or below epigastrium
- Localized to a point
 Pricking or piercing or
- stabbing type of pain
 Precipitated by movement of
- neck or arms or respiration

Any effort related pain fitting in previous category, relieved by rest or NTG in 1-2 min

STABLE ANGINA: GENERAL MANAGEMENT

1. Manage factors potentaiting angina

- Anemia, Thyrotoxicosis, Pregnancy, febrille illness
- Hypertension, Ventricular hypertrophy, CHF
- Tachy or brady-arrhythmia
- Drugs : bronchodilators, steroids

2. Risk factor control

- 3. Other atherosclerotic CV disease : PVD, stroke
- 4. Secondary prevention : Statins, BB, ACE-I

ESSENTIAL INVESTIGATIONS

- 1. Hemogram
- 2. Urea, Creatinine, Electrolytes
- 3. Sugar, HbA1C
- 4. Lipids
- 5. Liver function test
- 6. ECG
- 7. Plain X-ray chest

INVESTIGATIONS

MANAGEMENT

DESIRABLE INVESTIGATIONS

- 1. Echocardiography
- 2. Exercise Treadmill Test
- 3. Thyroid Function Test
- 4. Iron profile

2. Echocardiography for LV function or structural heart

patient is ambulatory and ECG is interpretable

3. Risk stratify by exercise treadmill test in low, intermediate

or high risk (DUKE risk score) for cardio-vascular events, if

Angina uncontrolled on optimal medical therapy

Non-ambulatory patient or un-interpretable ECG

High risk on exercise stress test for possible

5. Uric acid

OPTIONAL INVESTIGATIONS

- 1. Stress radionuclide/ echocardiographic imaging
- 2. CT scan including multi-slice coronary angiography
- 3. Coronary Angiography
- 4. Coronary Fractional Flow Reserve
- 5. Intra-vascular Ultrasound/OCT

MANAGEMENT AT PHC/ CHC LEVEL

MANAGEMENT AT DISTRICT HOSPITAL LEVEL

Echo reveals abnormality

re-vascularization

1. Optimise anti-anginal treatment

4. Refer to tertiary centres if:

MANAGEMENT AT TERTIARY LEVEL

 Reassess and optimise drug therapy: If uncontrolled choose from trimetazidine, nicorandil ranolazine and ivabid
 Risk stratify with exercise treadmill test if not already done
 Stress imaging if following:

 Non ambulatory patient
 Abnormal or uninterpretable baseline ECG
 Exercise treadmill test result is equivocal
 Compromised LV function

ATIONS

- 1. Control angina : Metoprolol Add nitrates if symptoms not controlled
- 2. ECG for Q waves, ST T changes, BBB or chamber enlargement
- 3. Aspirin & high intensity statins
- 4. Refer to higher centre electively

RISK CATEGORIZATION

A. Very high:

disease

Based on clinical features, GRACE score & TIMI score

- -Acute LVF -Hypotension
- -Uncontrolled Ventricular arrhythmia -Severe MR
- **RISK CATEGORY MANAGEMENT**

Low/ Intermediate Risk Group

- 1. Optimal anti-anginal therapy
- 2. Follow up 3-6 monthly at primary/ secondary care centre
- 3. Refer to tertiary centre when change in symptomatic status

High Risk Group

- 1. Discuss pros and cons of possible revascularization and dual anti-platelet therapy
- 2. Angiography, if any of following
 - Angina not controlled on optimal medical therapy
 - High risk on non-invasive testing
- Cardiac arrest survivor or documented VT

- B. High Risk:
- -GRACE score > 140 or TIMI score >4 C. Intermediate Risk:
- -GRACE score 109-140 or TIMI score 2-3 D. Low Risk:
 - -Grace score <108 or TIMI score 0-1

REVASCULARIZATION

- 1. Revascularize if anatomy is suitable
- 2. Prefer CABG over PCI in DM with multivessel disease or left main disease
- 3. Complete re-vascularization is preferable
- 4. Use invasive functional and imaging modalities (FFR, IVUS, OCT) when indicated
- 5. Stress on continuing dual anti-platelets (aspirin and clopidogrel) after PCI

DRUGS & DOSAGE

Anti-platelets

Aspirin 75 mg OD
 Clopidogrel 75 mg OD (if intolerant to aspirin)

Statins:

Atorvastatin: 40-80 mg OD Rosuvastatin: 20-40 mg OD

Ace-inhibitor

Ramipril: 2.5-10 mg OD Enalapril: 2.5-10 mg BD

Anti-ischemic:

1. Metoprolol: Short acting: 25-100 mg BD Long acting: 25 -100 mg OD

2. Nitrates:

Isosorbide mono-nitare: 20 to 60 mg in 2 devided dose Nitroglycerine sustained release: 2.6 to 6.5 mg BD

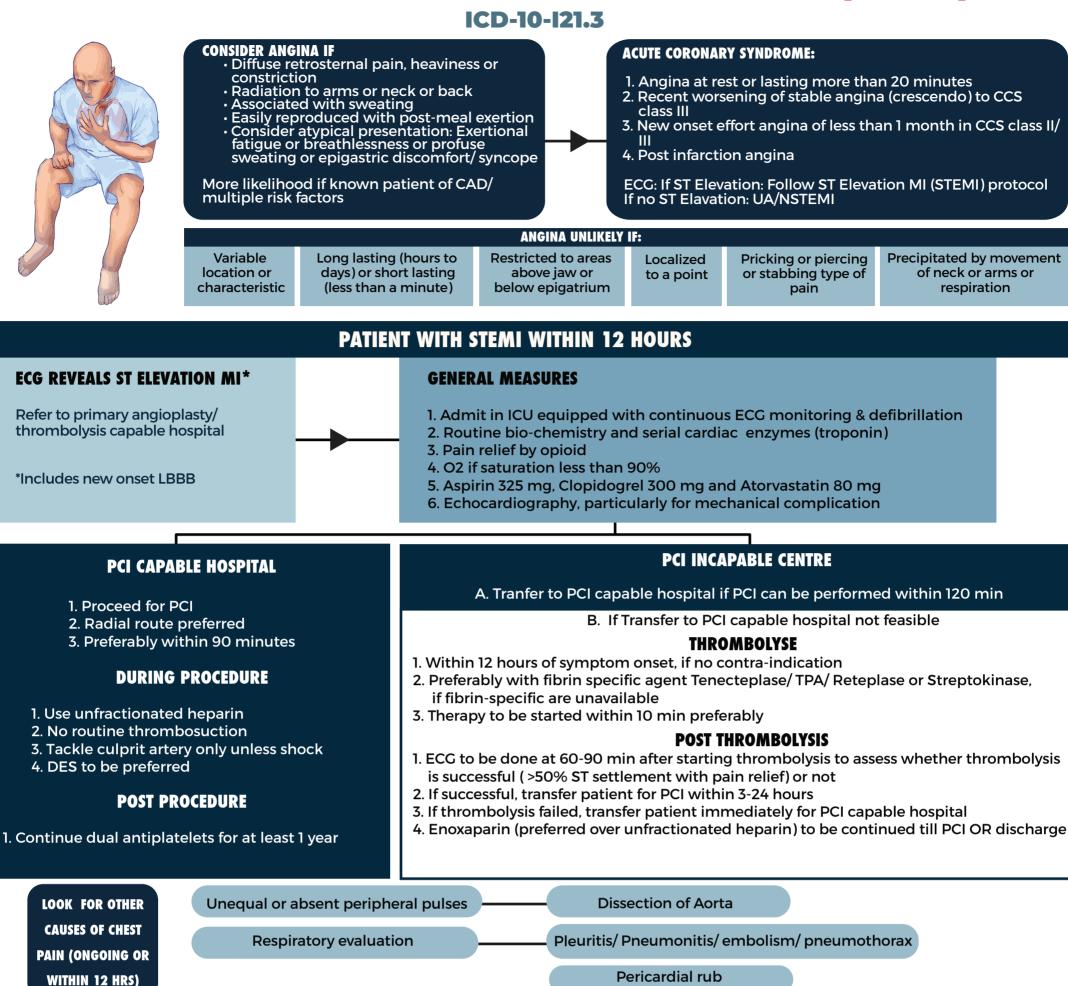
- 3. Calcium channel blockers: Verapamil 40-80 mg TDS Diltiazem 30 to 90 mg TDS
 - Nicorandil 5-10 mg BD
- 4. Nicorandil: 5-10 mg BD
- 5. Ranolazine: 500 -1000 mg BD
- 6. Trimetazidine: 20 mg mg TDS

• KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES : STRENGTHEN SECONDARY PREVENTION WITH STATINS, BB & ACE-I





Standard Treatment Workflow (STW) for the Management of **ST ELEVATION MYOCARDIAL INFARCTION (STEMI)**



Neuralgia or herpes

PATIENT WITH STEMI IN 12-24 HOURS

Transfer to PCI capable hospital immediately					If ongoing pain, thrombolysis and transfer immediately				
PATIENT WITH STEMI AFTER 24 HOURS									
F	Angiography with	a view to PCI	only if any	of follo	wing/ Contra indica	tions of angio	graphy:		
Recurrent anginal pain not controlled by medical therapy	rdiogenic shock	Acut	te LVF		Mecahnical complication	Dynam chai		ver	nreatening ntricular nythmias
	ABSO	UTE CONTRA-	INDICATIO	NS TO T	HROMBOLYIC THER	APY:			
hemorrhage or stroke of		neoplasm or nalformation	Recent (w month) r trauma/ s head in	major urgey/	Recent (within 1 month) major GI bleed	Known blee tendency (e) menstrual b	cept Ao	rtic dissection	Severe uncontrolled hypertension
	DRU	GS & DOSA	GE				S	TEMI DIAG	NOSIS*
Anti-platelets 1. Aspirin: Loading dose 2. Clopidogrel: Loading do 3. Prasugrel: Loading do 4. Ticagralor: Loading do Anti-ischemic: Metoprolol: Short acting: 25-100 Long acting: 25-100 Nitroglycerine susta Nitroglyceri	dose 300 mg follo ose 60 mg follower ose 180 mg follower 0 mg BD 0 mg OD itare 20 to 60 mg ained release 2.6 to 25 mcg/ min infus 80 mg OD	nutes	D BD	1. Un 60 fol inf 50 2. Er TH Tene 3. 4 4 8 Rete 10 r Alte 11 0 5 Stre	Anti thrombotics: Ifractionated hepari U/Kg (maximum 50 Ilowed by 12 U/Kg ho fusion to maintain A 0-70 sec noxaparin: 1 mg/Kg S hrombolyic Therap ecteplase 5 mg IV bolus if 60-1 0 mg IV bolus if 70-8 5 mg IV bolus if 70-8 5 mg IV bolus if 70-8 5 mg IV bolus follow 0.75 mg/Kg over 30 r 50 Kg weight, then 0 0 ver 60 min up to 35 ptokinase 5 million units IV ove	000 U) burly APTT at C 12 hrly y: 70 Kg 80 Kg re than at after 30 red by min upto 0.5 mg/Kg mg	Preferably <60 mins Primary Rescue No Yes Preferably Corona angiogra	PCI PCI Preferably «90mins («60 mins) («60 mins) presenters PCI Successful fibrinolysis Impresenters * The tir confirme ECG ides the First	EMS or non primary-PCI capable centre PCI possible <120 mins? Yes No Preferably within 30 mins Preferably within 30 mins Preferably within 30 mins

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



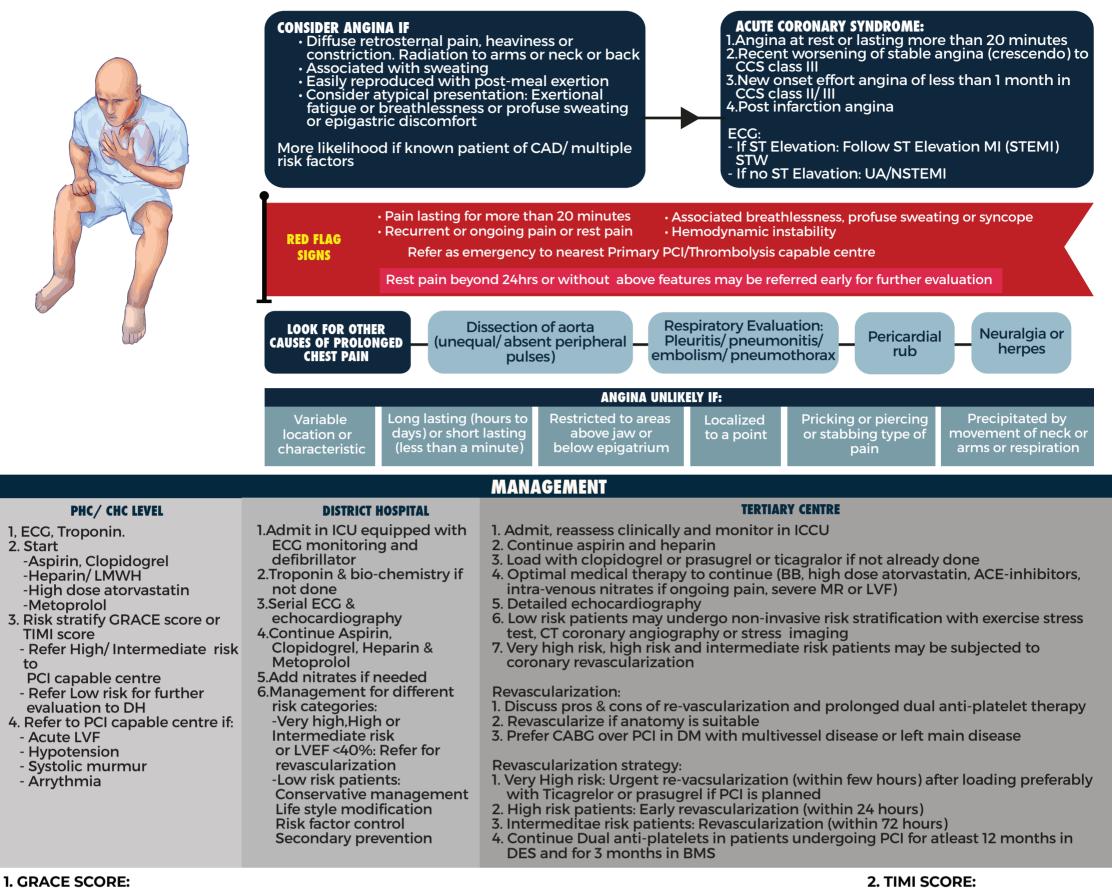




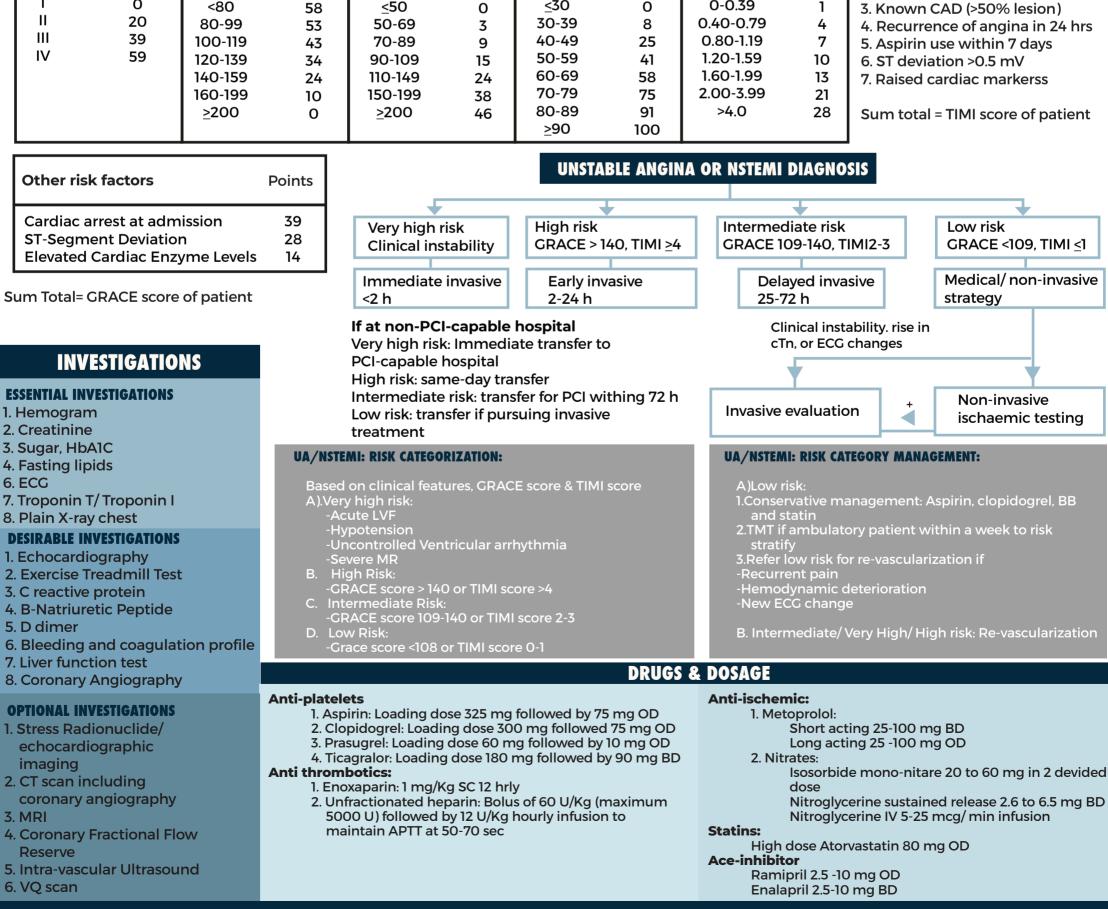
One point for each of following

Standard Treatment Workflow (STW) for the Management of **UNSTABLE ANGINA/ NSTEMI**

ICD-10-120.0



		_								2. 1101 3001(2.
Killip Class	Points	SBP1 mm Hg	Points	Heart rate Beats/ min	Points	Age. y	Points	Creatinine Level, mg/ dL	Points	One point for each of follo 1. Age >65 yrs
1	0	<80	58	≤50	0	<u><</u> 30	0	0-0.39	1	2. More than 3 risk factors 3. Known CAD (>50% lesion



🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURE

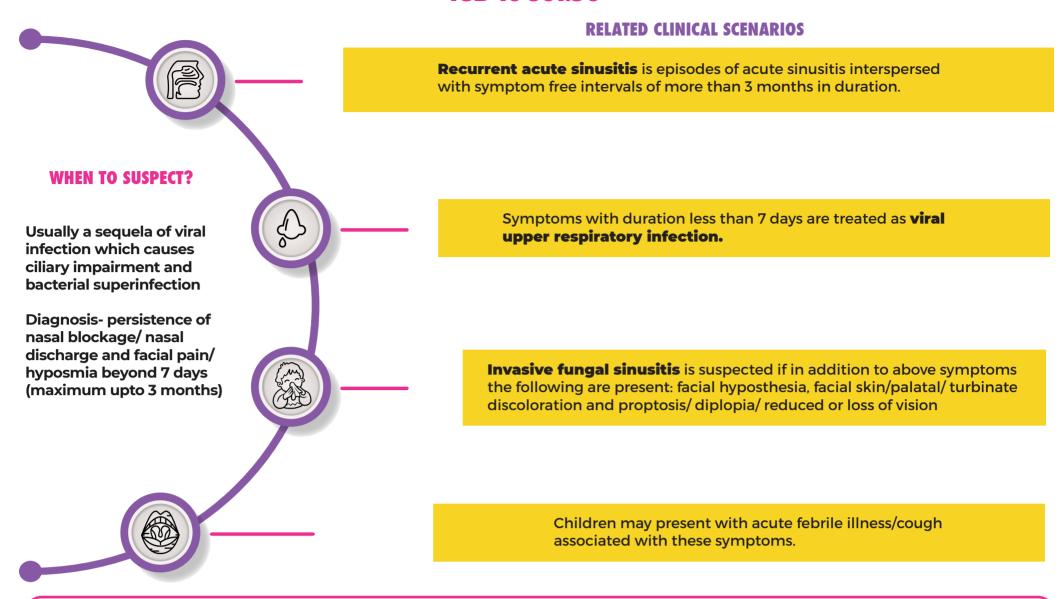


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Standard Treatment Workflow (STW) for the Management of **ACUTE RHINOSINUSITIS** ICD 10 J01.90



ALTERNATIVE CLINICAL SCENARIOS

- Consider alternate diagnosis if: Unilateral symptoms/ Bleeding/ Crusting/ Cacosmia (foul smell)
- Rule out other contributory factors: Allergy/ upper alveolar dental caries/ DNS/ LPR/ smoking.
- Rhinorrhoea and nasal congestion in second trimester of pregnancy is considered hormonal in etiology and is to be managed with saline irrigation/ drops

RED FLAGS FOR REFERRAL TO DISTRICT HOSPITAL

- Known diabetic/immunocompromised
- Suspicion of complications viz. (A) Orbital involvement (Periorbital edema/ erythema, displaced globe, ophthalmoplegia, visual disturbance); (B) Meningitis/ altered sensorium; (C) Frontal fullness.
- Non-resolution with oral antibiotics for ten days
- · Pointers of invasive fungal sinusitis (Facial hypoesthesia, facial skin/palatal/turbinate discoloration)

PRELIMINARY

- Anterior rhinoscopy: Discharge, bleeding, crusting, polyposis
- Oral examination: Dental caries, post nasal drip, palatal discolouration
- Assess for contributory factors listed above

DESIRABLE

Nasal endoscopy

October/ 2019

Desirable in non-resolving/worsening cases despite antibiotic therapy

- Endoscopy- for guided nasal swabs/ KOH smear
- CT PNS (for suspected complications / non-resolving cases on antibiotics for 14 days)
- Screen for Diabetes / Immunodeficiency

MANAGEMENT

PHC / PRIMARY LEVEL

Duration of treatment 7-14 days

- Oral antibiotics- Amoxycillin/ Co-amoxyclav for 7-10 days. Levofloxacin and Azithromycin can be opted for patients intolerant/ sensitive to penicillins.
- Topical budesonide/ mometasone nasal spray once/twice a day for 2 weeks provides earlier symptomatic relief.
- Normal saline nasal washes help in clearing secretions and improved effect of topical medications
- · Topical/ oral decongestant (Oxymetazline/ pseudoephedrine) for 3-5 days relieves symptoms.
- Adequate hydration and steam inhation.
- · Antihistaminics (patients with co-existing allergy).

INDICATIONS OF PARENTERAL ANTIBIOTIC THERAPY

- Orbital/intracranial complications
- Non-resolution of symptoms with atleast 7 days of oral antibiotics
- Worsening of symptoms while on oral antibiotics

DISTRICT HOSPITAL

- Surgical interventions to manage: Underlying anatomical conditions causing recurrent acute sinusitis like- DNS/ adenoid hypertrophy/ anatomical variations seen on CT
- Ophthalmology referral for suspected intraorbital complications
- · Dental deferral for suspected dental origin infection.
- Invasive fungal sinusitis- start antifungal medications, control underlying immunocompromising co-morbidity and consider debridement.

TERTIARY LEVEL

Cases of acute invasive fungal sinusitis/ complicated acute bacterial sinusitis and patients with immunocompromised status may be referred for management.

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



CT: Computerized Tomogram **PHC:** Primary Health Center

DNS: Deviated Nasal Septum **LPR:** Laryngo Pharyngeal Reflux

REFERENCES

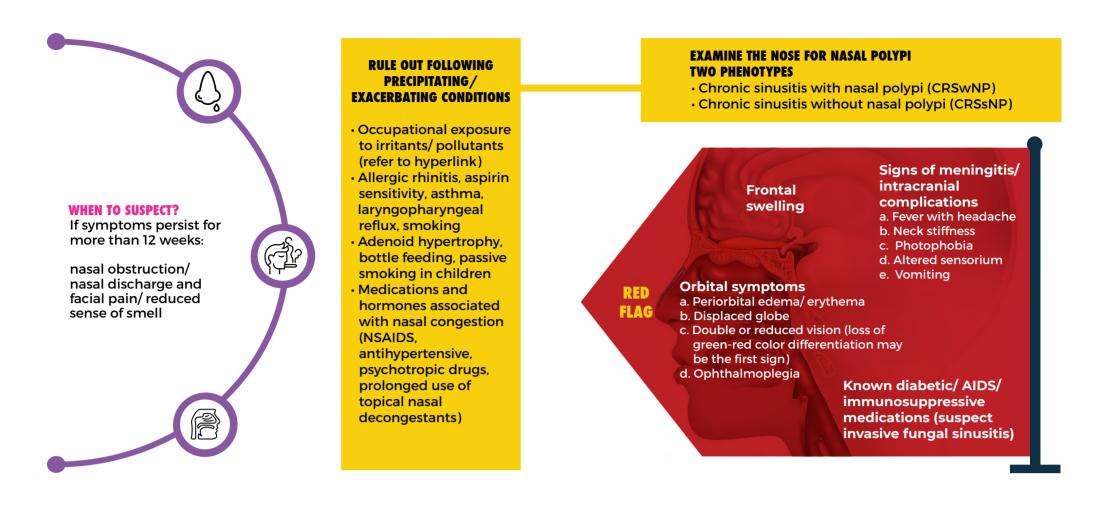
- 1. Indian Council of Medical Research. Treatment Guidelines for Antimicrobial Use in Common Syndromes. New Delhi, India, 2017.
- 2. Fokkens W, Lund V, Mullol J, et al. EPOS 2012: European Position Paper on Rhinosinusitis and Nasal Polyps 2012. Rhinol 2012;50(Suppl 23):1-298.
- 3. Sharma V, Saxena RK, Sharma S, Sharma G, Dhasmana DC, Mishra KC. Comparative Efficacy and safety of various anti-microbials in patients of acute rhinosinusitis at tertiary-care hospital in Uttarakhand. Indian Jour Otol Head & Neck Surg, 2011, Oct ; 63 (4): 364 9
- 4. Blomgren K, Eliander L, Hytönen M, Ylinen S, Laitio M, Virkkula P. How patients experience antral irrigation. Clin Med Insights Ear Nose Throat. 2015;8:13-7.



Department of Health Research Ministry of Health and Family Welfare, Government of India



Standard Treatment Workflow (STW) for the Management of CHRONIC RHINOSINUSITIS ICD 10 - J32.9



TREATMENT OF CRS

 Mild/ moderate symptoms (no significant congestion/ discharge/ polypi/complications)

- 1. Address etiology and exacerbating factors.
- 2. For allergic rhinitis, antihistamines and nasal steroid spray to be given.
- 3. Saline nasal wash
- 4. Steam inhalation

In presence of nasal purulent discharge

least 2weeks.

1. Culture directed antibiotics to be considered

2. If culture is negative, empirical antibiotics (Amoxycillin/

sinusitis which is to be treated with metronidazole

- 5. Stretching exercises and yoga are very effective for nasal congestion
- 6. Topical (oxymetazoline/ xylometazoline) and oral decongestants are associated with cardiovascular risks and rebound phenomenon. Hence, careful patient selection and short course treatment to be followed.
- 7. Intra nasal steroid sprays for 6-8weeks (Fluticasone proprionate/

IN ALL PATIENTS, ESPECIALLY IN THE PRESENCE OF NASAL POLYPI, RULE OUT ALLERGY/ALLERGIC RHINITIS

- 1. Consider allergen avoidance
- 2. Skin prick test
- 3. Co-existing bronchial asthma needs to be treated

Fluticasone furoate/ Mometasone) after discussing risk - benefit cost issues with patient regarding steroid sprays If no symptomatic relief to above treatment, perform nasal endoscopy and consider NCCT of paranasal sinuses

Co-amoxyclav/ Fluoroquinolone/ Roxithromycin) to be given for at

3. Upper dental (particularly 1st molar) infection may cause maxillary

HYPERLINK

(https://www.dovemed.c om/diseases-conditions/ airborne-irritant-induce d-sinusitis/)

• In the presence of nasal polypi, initial nasal steroid spray and subsequent endoscopic surgery is to be planned.

 Short course of oral steroid (Prednisolone 0.5 mg/kg for 5 - 10 days) provides temporary relief in nasal obstruction in extensive polypi.
 Steroid therapy is not a replacement for surgery.

Identification of precipitating or exacerbating factors is the key to successful treatment outcome. Always rule out DNS/ nasal polypi in CRS, as surgical treatment may be necessary for complete resolution of symptoms.

Ensure adherence to nasal saline washes / regular physical activity / medications. Educate patients on correct technique of using steroid nasal sprays and nasal irrigation. Prolonged use of topical nasal decongestant beyond 5-7 days may cause rebound congestion and rhinitis medicamentosa and to be strongly discouraged.

ABBREVIATIONS

CT: Computerized Tomogram

AIT: Allergen Immuno Therapy

DNS: Deviated Nasal Septum

REFERENCES

• Fokkens W, Lund V, Mullol J, et al. EPOS 2012: European Position Paper on Rhinosinusitis and Nasal Polyps 2012. Rhinol 2012;50(Suppl 23):1-298.

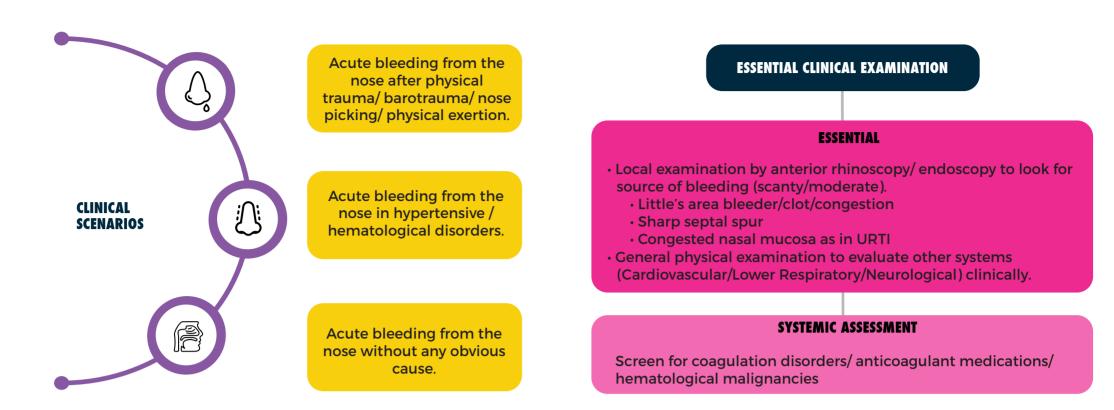
- Cain RB, Lal D. Update on the management of chronic rhinosinusitis. Infect Drug Resist. 2013;6:1-14.
- Ah-See KL, MacKenzie JM, As-See KW. Management of chronicrhinosinusitis. BMJ. 2012;345:e7054.

• Slovick A, Long J, Hopkins C: Updates in the management of chronic rhinosinusitis. Clin Pract. 2014;11(6):649-63. 10.2217/cpr.14.71



Ministry of Health and Family Welfare, Government of India

Standard Treatment Workflow (STW) for the Management of **EPISTAXIS** ICD-10-R04.0

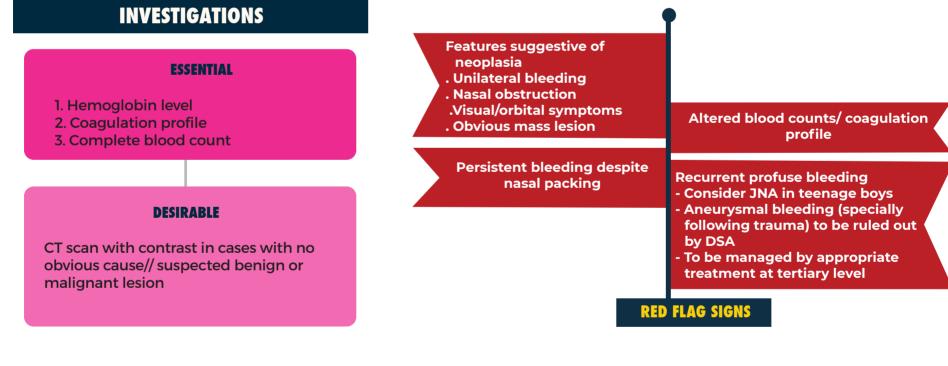


MANAGEMENT

STEP-WISE MANAGEMENT PRINCIPLE

- 1. Ensure patent airway/ avoid aspiration by head down/lateral positioning
- 2. Restore hemodynamic stability by intravenous fluid replacement/transfusion
- 3. Control bleeding/bleeder by
 - Bidigital compression of nose for 10 minutes in Trotter's position (cotton pledgets soaked in 4% xylocaine with adrenaline may be used)
 - Short term tab labetalol will take care of uncontrolled hypertension Chemical/electrocauterization of bleeder in Little's area
- 4. Tamponade of bleeders by anterior nasal packing/epistaxis balloon
- 5. Posterior nasal packing if bleeding is not controlled with above measures
- 6. Antibiotic prophylaxis and hospitalizarion is recommended after nasal packing
- 7. H2blockers/ PPI to be given in case of blood aspiration to avoid gastritis
- 8. Persisting bleeding despite nasal packing > consider arterial ligation (sphenopalatine / anterior ethmoidal artery).
- 9. Selective embolization is an alternative to surgery
- 10. Address identified etiology, if any





FOLLOW UP SERVICES

- 1. Continued nasal lubrication for 2 weeks with liquid paraffin
- 2. Repeat anterior rhinoscopy/ endoscopy to know/confim the cause of bleeding
- 3. Oral hematinics to be considered if needed

QUALITY ASSESSMENT PARAMETERS

- 1. Recurrence of episodes
- 2. Improvement in hemoglobin level over a period of time.

POINTS TO PONDER WHILE MANAGING EPISTAXIS

- 1. Epistaxis in children is almost always anterior and from Little's area, consequent to mucosal drying by dry air.
- 2. Epistaxis in adults is often related to hypertension and arises posteriorly from the posterior end of inferior turbinate
- 3. Initial non-invasive methods may suffice in a large majority of patients.

ABBREVIATIONS

JNA: Juvenile Nasopharyngeal Angiofibroma **DSA:** Digital Subtraction Angiography

CT: Computerized Tomograms **URTI:** Upper Respiratory Tract Infection

W KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



Department of Health Research Ministry of Health and Family Welfare, Government of India



Standard Treatment Workflow (STW) for the Management of HEARING IMPAIRMENT IN PEDIATRIC AGE GROUP (0 - 12 YEARS) ICD 10 H90.5

Disabling hearing impairment (31 or more dB HL in better ear) may affect language development and learning outcomes and hence needs urgent intervention

WHEN TO SUSPECT IN CHILDREN

- 1. Parental concern about delayed speech, language, and developmental delay (refer to red flags)
- 2. Family history of Hearing Loss (HL).
- 3. Exposure to ototoxic drugs/ hyperbilirubinemia requiring exchange transfusion/ Neonatal ICU stay for > 3 days.
- 4. In-utero infections (CMV/ rubella/ syphilis/ herpes/ toxoplasmosis)
- 5. Syndromes (NF) Or neurodegenerative disorders (Hunter syndrome, FA) associated with HL.
- 6. Post-natal infection known to cause HL (Meningitis) 7. Head Trauma
- 8. Recurrent/persistent (>/=3 months) middle ear disease
- 9. Chemo/ Radiotherapy to head and neck

EVALUATION

ESSENTIAL

- 1. Clinical examination to look for ear canal deformities, tympanic membrane and middle ear status by otoscopy/ otoendoscopy.
- 2. Age appropriate audiological/behavioral observation tests in a soundproof room by audiologist/ ENT specialist.
- 3. Tympanic membrane mobility test/ tympanometry.

RED FLAGS POINTING FOR URGENT HEARING EVALUATION

- 6months- no head turning to the side of calling
- · lyr- no babbling/speech like sound production
- 1.5yrs- not saying mama/papa/dada or other names
- \cdot 2yrs-not pointing to pictures/ body parts when named or speaking less than 10 words
- 3 yrs- does not understand action words or not asking for things by names or not speaking small sentences.
- At any age- has regressed or lost previously acquired speech/language milestones

MANAGEMENT

GUIDING PRINCIPLES



Middle ear fluid (OME) may be



- Community based hearing screening: i. May be co-ordinated with immunization schedule
 - ii. By primary health care workers. iii. Using calibrated noisemakers/ toys
- All children who fail preliminary screen to undergo detailed evaluation at health care facility.

COMMON CAUSES OF HL

1. Impacted wax

SNHL

- 2. Middle ear fluid assciated with adenoid hypertrophy/ cold climate
- 3. Tympanic membrane perforation
- 4. Sensorineural Hearing loss (SNHL) due to various causes as indicated earlier

vision by ENT specialist relieves hearing impairement Appropriate surgery is to be planned for tympanic membrane perforation For non surgical	ed with adenotonsillar disea needs to be treated. Initially I treatment and surgery to be ed for OME persisting for months/ earlier in the presence eech and language delay -surgical condidates/ delaye I management, amplification aring aid to be reinforced in bilateral CHL.	amplification, preferential seating in classroom e of Periodic evaluation for hearing aid users for mould fitting	Screening for developmental delay by pediatrician/ psychologist			
	DIVISION OF RESPONSI	BILITIES				
PHC LEVEL		DH LEVEL				
 Suspect HL Initial evaluation Referral if initial evaluation is suggestive of HL Follow up of rehabilitated/ treated patients with H Prevention of HL 	2. Hearing aid dis 3. Rehabilitation k 4. Appropriate su 5. Training progra	 Audiometric evaluation by Audiologist/Otolaryngologist Hearing aid dispensing (mould fitting and HA programming) Rehabilitation by speech therapist Appropriate surgery for CHL Training programme for parents of hearing impaired children to enhance pre-school language development 				
TERTIARY LEVEL		QUALITY ASSESSMENT PA	RAMETERS			
		 Short term: Quality of amplification 	ition using			

- Surgical intervention options : Cochlear implant / BAHA (as per ADIP quidelines)
- Interdisciplinary team based interventions in children with multiple disabilities.
- electroacoustic objective measures and culturally appropriate subjective questionnaire tools
- Long term (Desirable): Use CBR matrix based measurement for ensuring holistic rehabilitation

FOLLOW UP SERVICES

1. Home visits by Health Worker/ASHA to ensure utilization of assistive devices and support parents to enhance language development.

- 2. School visits to educate teachers and normally hearing children to include their peers with hearing disability in the school environment
- 3. Home/ school visit by social worker for evaluation of social/educational/livelihood/justice and empowerment domains of the child

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

ABBREVIATIONS

ADIP: Assistance to disabled persons for purchase/fitting of aids and appliances

BAHA: Bone Anchored Hearing Aid **CBR**: Community Based Rehabilitation **CMV**: Cyto Megalo Virus

FA: Friedreich Ataxia **NF**: NeuroFibromatosis **OME**: Otitis Media with Effusion

REFERENCES

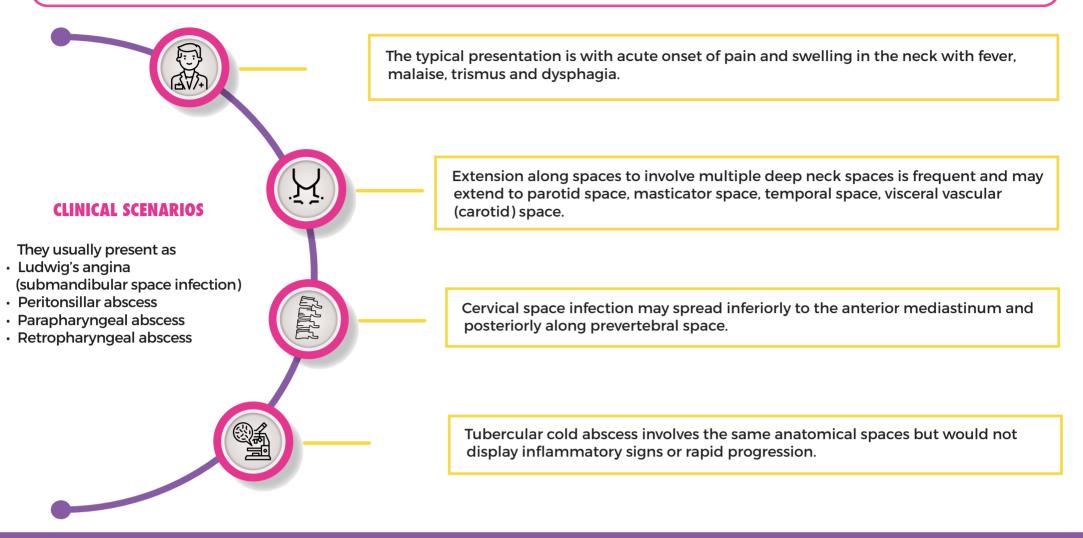
- Indian Council of Medical Research. Audiological evaluation protocols. Task force project on prevalence and etiology of hearing impairment, New Delhi. 2015
- Ramesh A, Jagdish C, Suman Rao PN et al. Low cost calibrated mechanical noisemaker for hearing screening in resource constrained settings. Indian Journal of Medical Research. 2012, 135: 170 176.
- Rathna.B.Shetty. Manual for training parents of hearing impaired children (Kannada : Kivudu makkalige kalisuva vidhana). Parents association of deaf children. Mysore.
- Chapal Mkhasnabis, Karen Heinicke Motsch (eds.) Towards community based inclusive development. World Health Organisation: 2010.
- Margaret Lavina Fernandes. Guidelines to establish a community based rehabilitation program for hearing impaired children in medically underserved areas. St. John's Medical Journal, 2018 (1), 5: 14 27
- ADIP Guidelines : http://disabilityaffairs.gov.in/content/page/adip-scheme.php





Standard Treatment Workflow (STW) for the Management of **NECK SPACE INFECTION** ICD-10-J36, J39.0, K 12.2, J39.1

Rapidly progressive bacterial infections which spread along facial planes and spaces of head and neck region. They may be fatal unless emergently treated. Most of these infections are secondary to dental infection.



SYSTEMIC ASSESSMENT

Screen for diabetes mellitus, HIV infection, agranulocytosis and immunosuppressive therapy or chemotherapy. Signs of inflammation may be less marked and disease course may be more rapidly progressive in immunocompromised patients.

CLINICAL EXAMINATION

- Airway assessment to rule out stridor or respiratory compromise
- Look for signs of dehydration
- Monitor temperature, heart rate, respiratory rate, BP, and signs of sepsis/ septic shock.
- Oral cavity examination to check jaw opening, condition of teeth and floor of mouth
- Oropharyngeal examination to check for inflammed medially displaced tonsil & uvula and bulge in lateral pharyngeal wall
- Palpation of neck for lymph nodes, cellulitis, abscess or
 subsutaneous grapitus

RED FLAGS FOR REFERRAL TO DISTRICT HOSPITAL

- Breathing difficulty
- Trismus
- Torticollis/ neck stiffness
- Subcutaneous crepitus and skin discolouration or blisters suggest necrotizing fibrofascitis.
- Toxaemia
- Lower cranial nerve palsy

subcutaneous crepitus

- Cranial nerve examination to rule out lower cranial nerve palsies
- Facial puffiness suggestive of venous thrombosis
- Mediastinal extension

INVESTIGATIONS

ESSENTIAL INVESTIGATIONS

- 1. Contrast enhanced CT scan of head and neck is the standard in evaluation of neck space infections. If CT Scan facility is not available, following should be done:-
 - a. Lateral x-ray neck: Prevertebral soft tissue thickening >7 mm at the level of C2 or > 2/3rd of the width of the vertebral body at C6 is highly suggestive of retropharyngeal abscess. It may also demonstrate foreign bodies, subcutaneous air, air fluid levels and erosion of vertebrae.
 b. Ultrasound neck can suggest abscess and guide aspiration attempts.
- 2. Blood: Total and differential leukocyte count, blood sugar, urea
- 3. Abscess Cultures with Gram stain to direct antimicrobial therapy. Anaerobic culture, when available.

MANAGEMENT **PHC/PRIMARY LEVEL DISTRICT HOSPITAL INDICATIONS FOR 1&D** 1. Cautiously assess the 1. Hospitalization: As an emergency for close watch and intensive Necrotizing fibrofascitis airway. If found management. Abscess formation compromised, do 2. Airway management: In progressive disease, in view of impending • No response to antibiotics over endotracheal intubation/ airway compromise, consider securing the airway early. During acute 48-72 hours consider tracheotomy respiratory difficulty, tracheostomy should be done if intubation is Deterioration despite antibiotics Immediately gain an IV difficult over 24 hours access for hydration, broad 3. Correction of fluid and electrolyte imbalance Airway compromise or impending spectrum antibiotics and 4. Antibiotics: Early and aggressive IV antibiotic therapy with a airway compromise pain killers. combination of Crystalline Penicillin, Aminoglycoside and Mediastinal spread 3. Transfer the patient to Metronidazole or Clindamycin is preferred. Vascular complication like venous hospital with facility for 5. Incision and drainage: Peritonsillar abscess is drained intraorally. All thrombosis surgical drainage other abscesses are drained via an external approach

QUALITY ASSESSMENT PARAMETERSFOLLOW UP SERVICESComplete resolution of infection and follow up to ensure no recurrence;
treatment of initial cause of infection in tooth or tonsil.Consider cold tonsillectomy for patients with history of multiple
episodes of tonsillar abscess

ABBREVIATIONS

CT – Computerized Tomography

MRI – Magnetic Resonance Imaging

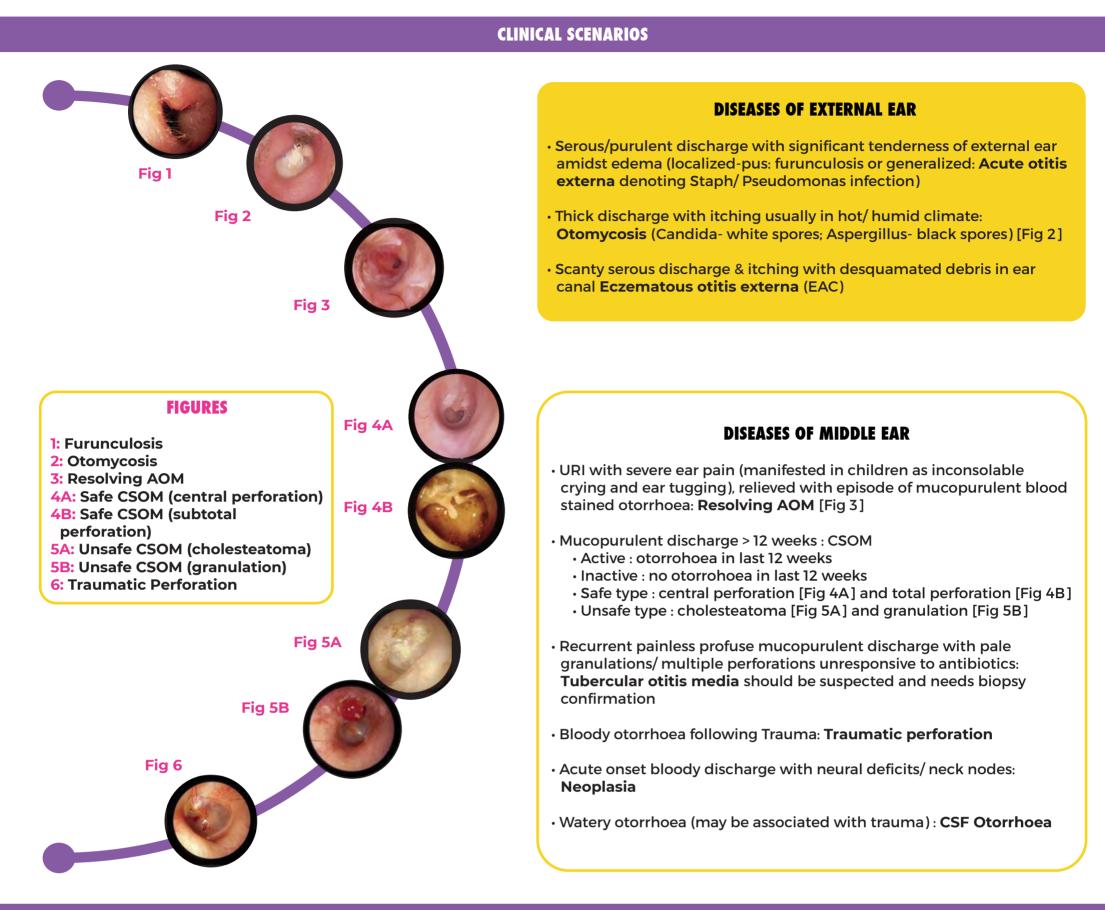
REFERENCES

- 1. Smith II JL, Hsu JM, Chang J (2006) Predicting deep neck space abscess using computed tomography. Am J Otolaryngol 27: 244-247.
- 2. Mayor GP, Millán JMS, Martínez VA (2001) Is conservative treatment of deep neck space infections appropriate? Head And Neck 23: 126-133.
- 3. Bottin R, Marioni G, Rinaldi R, Boninsegna M, Salvadori L, et al. (2003) Deep neck infection: A present day complication. A retrospective review of 83 cases. Eur Arch Otorhinolaryngol 260: 576-579.





Standard Treatment Workflow (STW) for the Management of OTORRHOEA ICD-10-H92.10



CLINICAL EXAMINATIONS

- Otoscopy as a part of Complete ENT examination by primary physician (Tele-otoscopy interpreted by physician)
- Hearing evaluation by conversation/ whisper/ Tuning forks tests
- General and systemic clinical examination

INVESTIGATIONS

- Pure tone audiometry
- Routine hemogram including blood sugar (fasting and postprandial)
- CT/ MRI in suspected complications (refer to red flags)
- Soft tissue x ray nasopharynx (To examine adenoid enlargement in children)
- Culture & sensitivity of aural secretions.

RED FLAGS FOR REFERRAL TO DISTRICT LEVEL

- Periaural abscess or cellulitis
- High grade fever, dizziness and toxic appearance
- Severe headache with neck stiffness/ vomiting / altered sensorium.
- Facial palsy/ Neurological defecits
- Diabetic with severe deep seated ear pain / neural defecits (Skull base osteomyelitis)
- Physical trauma with bloody/ watery discharge (suspected CSF leak)
- Suspected tuberculosis/ neoplasm

MANAGEMENT

PHC / PRIMARY LEVEL

- Acute otitis externa: Oral Ciprofloxacin/ Amoxycillin clavulanic acid combination for 7-10 days (2 weeks maximum) and analgesics. Ichthammol gycerine (1:9) packing of EAC in moderate to severe edema. Refer pus pointing furuncle to DH
- Otomycosis: Cleaning and Clotrimazole ear drops
- · Eczematous otitis externa: Ciprofloxacin ear drops with steroid combination.
- AOM / Resolving AOM: Oral amoxicillin / Erythromycin / Clarithromycin for 10 days. With no response in 3 days start Amoxycillin clavulanic acid combination for 10 days. Refer to DH if no resolution
- Inactive CSOM: Referral to DH for surgery.
- Active CSOM: Ciprofloxacin ear drops with dry mopping & referral to DH for surgery. A course of oral antibiotics maybe prescribed in ase of persistant otorrhoea after topical antibiotics
- Traumatic perforation: Topical antibiotics for otorrhoea if any and maintain ear dry till healing complete
- In case of suspicion of complications start intravenous Amoxycillin clavulanic acid combination and refer to DH

DISTRICT HOSPITAL

- Surgical interventions except neurosurgical interventions (eg I&D, tympanoplasty, mastoidectomy)
- Biopsy in suspected neoplasm
- Medical management of medical co-morbidities such as diabetes, tuberculosis, meningismus/ meningitis

TERTIARY LEVEL

Surgical management particularly of intracranial complications including neurosurgical interventions

- Patient to be educated for proper technique of ear mopping, contralateral lie (10 min) following instillation of drops & avoiding water entry e.g ear-plugs during bathing
- To ensure adequate immunization (measles/ H.Influenza/ Pneumococcus) in recurrent AOM and to adopt correct posture during breastfeeding while avoiding bottle feeding
- Pus culture sensitivity to guide antibiotic regime in recurrent/ complicated cases Patient education to refrain from indigenous (oil/ hot water/ acid etc) ear treatments

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

ABBREVIATIONS

CT: Computerized Tomogram **MRI:** Magnetic Resonance Imaging **AOM:** Acute Otitis Media **CSOM:** Chronic Suppurative Otitis Media **EAC:** External Auditory Canal **URI:** Upper Respiratory Infection

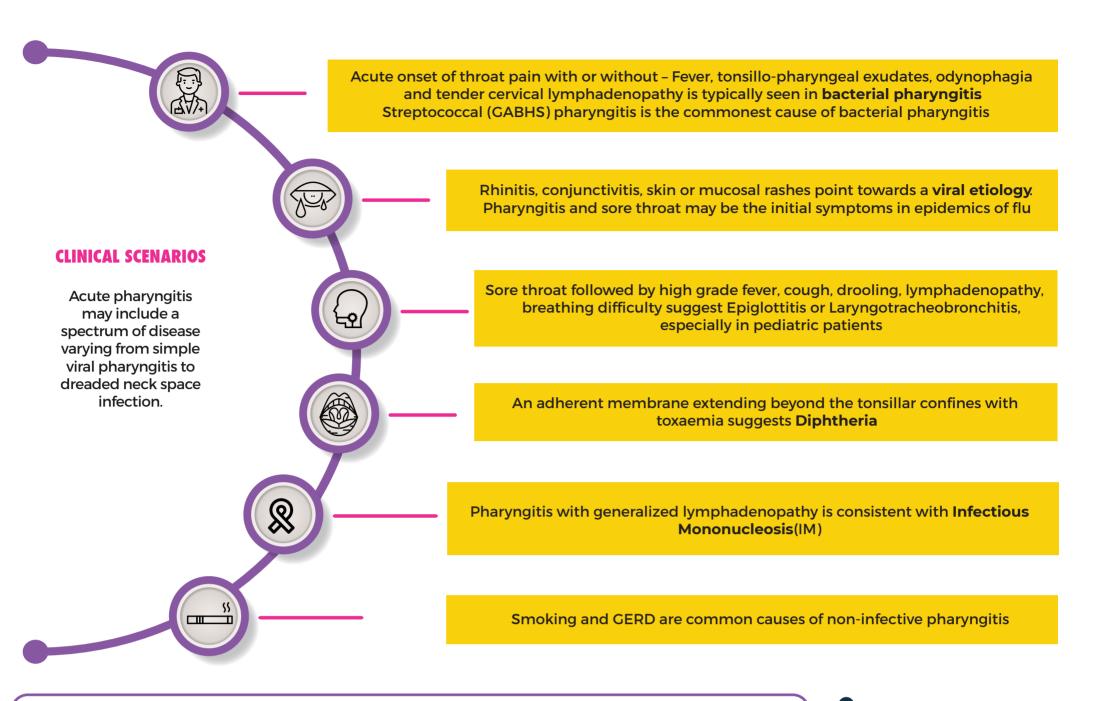
REFERENCES

- Otitis media (acute): antimicrobial prescribing. NICE guideline. Published: 28 March 2018. nice.org.uk/guidance/ng91
- Primary ear and hearing care training resource, Student's workbook: intermediate level. Chronic disease prevention and management. WHO 2006
- Primary ear and hearing care training resource, advanced level. Chronic disease prevention and management. WHO 2006
- Treatment Guidelines for antimicrobial use in common syndromes. ICMR. Department of Health Research. 2017
- Sagar P, Thakar A, Samant S. Otorhinolaryngology. In Paul VK, Bagga A. eds. Ghai Essential Pediatrics, 9th ed. New Delhi: CBS Publishers & Distributors; 2019. p.357-370



Ministry of Health and Family Welfare, Government of India

Standard Treatment Workflow (STW) for the Management of **PHARYNGITIS AND SORE THROAT** ICD-10-J02



CLINICAL EXAMINATION

PRELIMINARY

- Temperature chart: fever is usually absent or low-grade in viral pharyngitis
- · Check for vitals/ signs of dehydration due to compromised oral intake due to odynophagia
- · Complete oral and oropharyngeal examination with tongue depressor
- Palpate for cervical and generalized lymphadenopathy
- Rheumatic fever and acute glomerulonephritis are potential systemic complications of streptococcal pharyngitis
- · Hepatosplenomegaly can be found in IM
- A sandpapery scarlatiniform rash may be seen in GABHS infection whereas maculopapular rashes are seen with various viral infections and with IM empirically treated with penicillin

DESIRABLE

Assess Centor criteria and ascertain its score

RED FLAGS

- Generalized lymphadenopathy
- · Cardiac murmurs Purulent
- productive cough with tachypnea
- suggestive of LRTI Hot potato voice
 - Unilateral tonsillar

CLINICAL FEATURES	CENTOR SCORE	UNLIKELY TO HAVE GABHS	LIKELY TO HAVE GABHS	REQUIRE LAB TESTS TO CONFIRM GABHS INFECTION	• Ton me bey
Fever	1				COL
Anterior cervical lymphadenopathy	1	Score = 0-1	Score = 4	Score = 2-3	• Agr • Epi
Tonsillar exudate	1				
Absence of cough	1				

emargement
• Tonsillar
membrane going
beyond its
confines
Agranulocyosis

•	Ep	ide	em	ic c	of flu

INVESTIGATIONS									
ESSENTIAL	OPTIONAL	DESIRABLE							
Throat swab for culture, routine hemogram including total and differential leukocyte counts and peripheral smear to look for atypical lymphocytes (seen in IM).	GABHS rapid antigen detection test (RADT)	Lab tests to rule out EB Virus, Coxsackie virus, Herpes virus, fungal or Gonococcal pharyngitis							

MANAGEMENT						
PHC / PRIMARY LEVEL	DISTRICT HOSPITAL					
 Assess the patient for signs of toxicity, epiglottitis or oropharyngeal abscess Ensure vitals/ hydration of the patient Saltwater gargle, warm liquids, and rest may be helpful in relieving symptoms Ibuprofen or Paracetamol is recommended for analgesia Antibiotic therapy: a. Patients positive for all 4 Centor criteria to be treated with antibiotics without waiting for antigen testing or cultures b. Patients with Centor score of 2&3 to be treated with antibiotics only if antigen testing or throat swab culture is positive c. Patients with Centor score of only 1 not to be treated with antibiotics d. Amoxicillin (50 mg/kg/d in 2-3 doses orally) for 10 days is the first choice for GABHS infection. For patients who are sensitive for penicillin group, Erythromycin or Azithromycin is the antibiotic of choice Parenteral antibiotics (Ceftriaxone/ cefotaxime) and steroids are to be started when the airway is compromised due to suspected epiglottis/Croup. 	Management of complication e.g. • Deep neck space infection • Diphtheria • Epiglottitis • Croup					

FOLLOW UP SERVICES

Recurrent (more than 7 episodes in previuos year or 5/year in last two years or 3/year in last 3 years) tonsillitis episodes need to be evaluated for tonsillectomy.

🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

ABBREVIATIONS

GABHS: Group A Beta Hemolyticus Streptococcus **GERD:** Gastro Esophageal Reflux Disease LRTI: Lower Respiratory Tract Infection

EB: Epstein Barr **RADT:** Rapid Antigen Detection Test

REFERENCES

1. Shaikh N, Swaminathan N, Hooper EG (2012) Accuracy and precision of the signs and symptoms of streptococcal pharyngitis in children: a systematic review. J Pediatr. 160 (3): 487-493.

- 2. Centor RM, Allison JJ, Cohen SJ (2007) Pharyngitis management: defining the controversy. J Gen Intern Med. 22(1): 127-130.
- 3. Altamimi S, Khalil A, Khalaiwi KA, Milner R, Pusic MV, AI Othman MA (2009) Short versus standard duration antibiotic therapy for acute streptococcal pharyngitis in children. Cochrane Database Syst Rev. 21. CD004872.

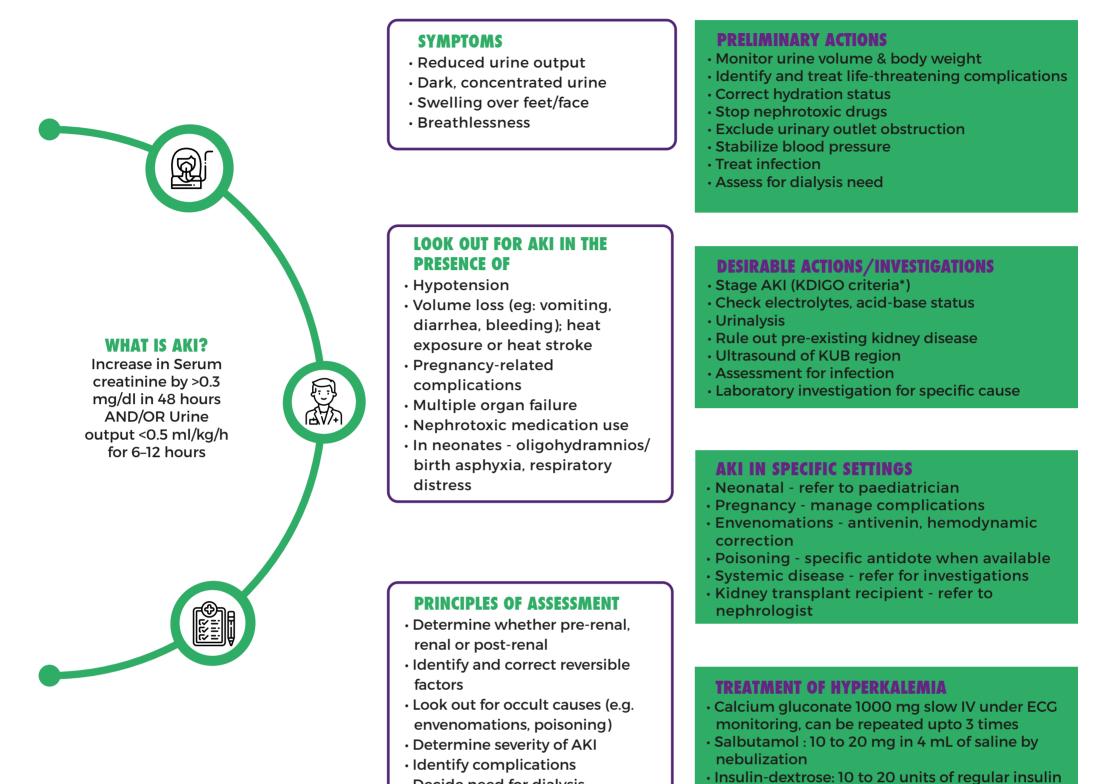


NEPHROLOGY





Standard Treatment Workflow (STW) for the Management of **ACUTE KIDNEY INJURY** ICD-10-N17.9



MANAGEMENT

PRIMARY CARE

October/ 2019

- Detailed history and physical examination
- Identify and correct volume deficit
- Stop nephrotoxic agents
- Identify and correct bladder outlet obstruction
- Give anti-snake venom if indicated
- Identify hyperkalemia and start treatment
- Identify pulmonary edema- start intravenous furosemide and oxygen
- PD if indicated
- Timely referral after stabilisation

RED FLAGS FOR URGENT REFERRAL

- Indications for dialysis
- Unexplained AKI
- Involvement of other organs
- Sepsis
- Systemic disease
- Complicated pregnancy

SECONDARY CARE

- Detailed history and physical examination
- Identify and correct volume deficit
- Stop nephrotoxic agents
- Identify and treat hyperkalemia, metabolic acidosis and pulmonary edema
- Identify and correct urinary tract obstruction (USG, CT)
- Detailed investigation for infections
- Manage pregnancy complications deliver if indicated
- Look for underlying CKD
- \cdot Dialysis (PD or HD)

INDICATIONS FOR DIALYSIS

- Fluid overload
- Pericarditis
- Hyperkalemia
- Severe metabolic acidosis
- Encephalopathy
- Severe uraemia
- To create space for fluids or blood products

TERTIARY CARE

- Detailed history and physical examination
- Identify and correct volume deficit

in 100 ml 25% or 50% dextrose

- Stop nephrotoxic agents
- Identify and correct urinary tract obstruction (USG, CT scan)
- Identify and treat hyperkalemia, metabolic acidosis and pulmonary oedema
- Detailed investigation for infections
- $\boldsymbol{\cdot} \text{ Manage pregnancy complications- deliver if indicated}$
- Look for underlying CKD
- Investigations for specific cause (including imaging, genetic tests)
- Kidney biopsy
- Dialysis (PD or HD)

FOLLOW-UP OF AKI

- UO > 1L, stable or falling creatinine, no symptoms: stop dialysis
- Not resolving for >2 weeks: CECT to exclude cortical necrosis; kidney biopsy as indicated
- Look for systemic diseases (e.g. vasculitis, myeloma, TMA)
- Serum creatinine and urine protein q 6-12 months for life

ABBREVIATIONS

AKI: Acute Kidney Injury **CECT:** Contrast-enhanced CT scan **PD:** Peritoneal dialysis **TMA:** Thrombitic microangiopathy

CKD: Chronic Kidney Disease **HD:** Hemodialysis

UO: Urine output **USG:** Ultrasonography

REFERENCE

***KIDNEY DISEASE:** Improving Global Outcomes (KDIGO) Acute Kidney Injury Work Group. KDIGO Clinical Practice Guideline for Acute Kidney Injury. Kidney Int, Suppl. 2012; 2: 1–138

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



Department of Health Research Ministry of Health and Family Welfare, Government of India



Standard Treatment Workflow (STW) for the Management of CHRONIC KIDNEY DISEASE (CKD) ICD-10-N18.3

WHEN TO LOOK FOR CKD

- History of long-standing nocturia, or constitutional symptoms
- Edema, hematuria, proteinuria or renal stones
- Long-term intake of painkillers or herbal medicines
- Family history of kidney disease
- Growth retardation, rickets, or proximal myopathy
- Unexplained hypertension or anemia
- · Longstanding diabetes, hypertension, CVD, stroke, PVD
- Systemic diseases (e.g. connective tissue disease)

EVALUATION OF NEWLY DIAGNOSED PATIENT WITH CKD

- Serum creatinine, electrolytes, bicarbonate
- Estimate glomerular filtration rate using CKD-EPI equation
- Urinalysis (examine sediment, proteinuria quantitation)
- Ultrasound of kidneys and urinary tract
- Calcium, phosphate, alkaline phoshatase, albumin
- CBC including peripheral blood film
- · Iron profile Serum iron, TIBC, TSAT
- HBsAg, anti-HCV

INITIAL ASSESSMENT FOR

- Confirmation of CKD diagnosis (repeat tests after 3 months)
- Staging and progression rate
- · Establishing cause of kidney disease
- Identify and treat reversible factors (hypertension, volume loss, obstruction, infection)
- · Look for complications (anemia, bone disease,
- dyselectrolytemias, CVD)

LIFESTYLE MEASURES FOR ALL CKD PATIENTS:

- Weight control/ weight gain monitoring in children
- Regular physical activity
- Reduce dietary salt intake to < 5 g/day
- Stop tobacco use in all forms
- Stop/moderate alcohol use
- Stop using unproven health supplements
- Do not use NSAIDS
- Avoid untested indigenous medicines

BP CONTROL (TARGET <130/80, 120/80 IF PROTEINURIA)

- Restrict dietary salt to < 5 g/day
- Use any anti-HT available in local pharmacy
- Diuretics eGFR > 45 : thiazide, <45 ml/min: furosemide; <30 ml/min: do not use potassium sparing agents
- ACEI/ARB preferred* for proteinuric patients (> 1 g/d)
- *caution/do not use if eGFR <30 ml/min, or Potassium >5.5 mEq/L

VACCINATION SCHEDULE FOR NEWLY DIAGNOSED CKD PATIENT

- If HBV -ve: 20 µg IM in each deltoid at 0,1,2 and 6 months
- In children complete primary vaccination schedule

ANEMIA MANAGEMENT

- Establish iron replete state
- If not iron replete, give oral iron
- Consider IV iron for dialysis patients
 and those not tolerating orally
- If Hb still <8 g/dl start erythropoietin, titrate to Hb 10-11 g/dl

MANAGEMENT OF HYPERPHOSPHATEMIA (PO4>5.5)

- Start with Ca-containing binders
- Non Ca-binders can be used if serum Ca >9 mg/dl, vascular calcification or low iPTH

DIABETES CONTROL (TARGET HBA1C <7%) Do not use metformin if eFGR <30

WHAT IS CKD?

Abnormalities of kidney structure or function, present for >3 months, with implications for health

NUTRITION

- Salt restriction < 5g/d. Protein 0.6-0.8 g/kg/day.
- DO NOT restrict proteins unless documented high protein user (dairy, white meat are good protein sources, mix different types of dal).

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- Restrict green leafy vegetables if eGFR <30 ml/min
- $\boldsymbol{\cdot}$ Avoid fruit juices, coconut water and carbonated beverages
- For children: ensure adequate protein intake appropriate for age.

LOW POTASSIUM FRUITS/ VEGETABLES:

Apple, pineapple, papaya, pear, tangerine, watermelon, grape, plum, cabbage, carrot, cauliflower, onion, radish, peppers, chillies, brinjal, cucumber, green beans, peas, rice, bread

VITAMIN D THERAPY

- Supplement 60,000 units cholecalciferol q2W
- Correction of acidosis with oral sodium bicarbonate
- Activated vitamin D if hyperparathyroidism

MANAGEMENT

PRIMARY CARE

- $\boldsymbol{\cdot}$ Detailed history and physical examination
- Identify and correct reversible factors
- Stop nephrotoxic agents
- Referral after stabilization

ADMISSION CRITERIA

- Initial evaluation or when patient presents with specific problems – like acute worsening, development of a new complication
- For creation of vascular access
- For PD catheter placement or initiation
- Initiation on HD and for kidney transplant

TERTIARY CARE

- Detailed history and physical examination
- Investigate to ascertain cause of CKD (imaging/biopsy/genetic studies)
- Tailor treatment to cause
- Identify and manage complications
- Vaccination
- Counseling: nutrition, lifestyle, pregnancy in women of child-bearing age
- Discussion regarding RRT
- Vascular access creation/PD catheter insertion
- Work-up for transplantation
- Send patient back to community with treatment plan

INDICATIONS FOR REFERRAL

- Initial evaluation of all newly diagnosed cases
- Rapid disease progression
- New complication
- Discussion for Renal Replacement Therapy (RRT)

DISTRICT HOSPITAL

- Detailed history and physical examination
- Investigate to ascertain cause of CKD
- Tailor treatment to cause
- Identify and manage complications
- Vaccination
- Identify and correct acute factors
- · Counseling: nutrition, lifestyle, pregnancy in women of child-bearing age
- Discussion regarding RRT
- Vascular access creation or PD Catheter insertion
- · Send patient back to community with treatment plan

PREPARATION FOR RENAL REPLACEMENT THERAPY

- eGFR < 30 : Preserve veins in the non-dominant arm for AV Fistula
- eGFR < 30 : discuss RRT options.
- eGFR < 15 : May need dialysis soon, counsel for AV fistula, list for transplant
- Dialysis start : depends on symptoms or eGFR <5 ml/min
- · Look for contraindications to HD or PD : discuss choice in those suitable for either

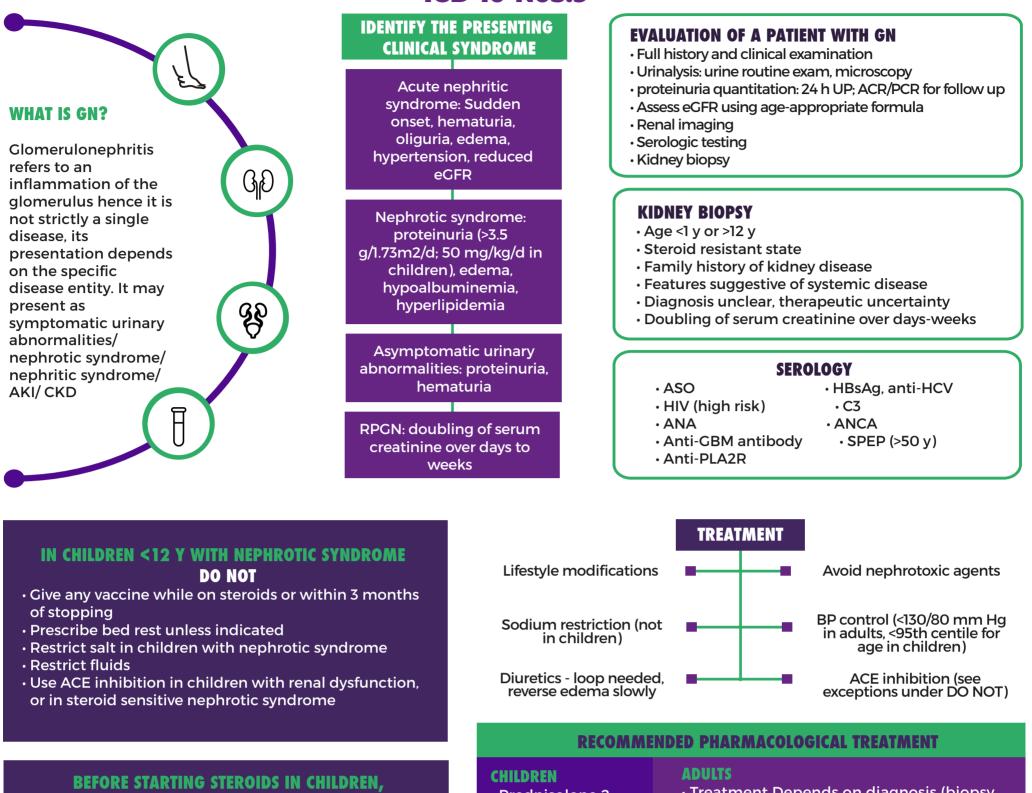
CONSERVATIVE CARE

If life expectancy limited, multiple comorbidities/personal preference
Decision-making should be shared with patient/family

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



Standard Treatment Workflow (STW) for the Management of **GLOMERULONEPHRITIS** ICD-10-N05.9



REMEMBER TO

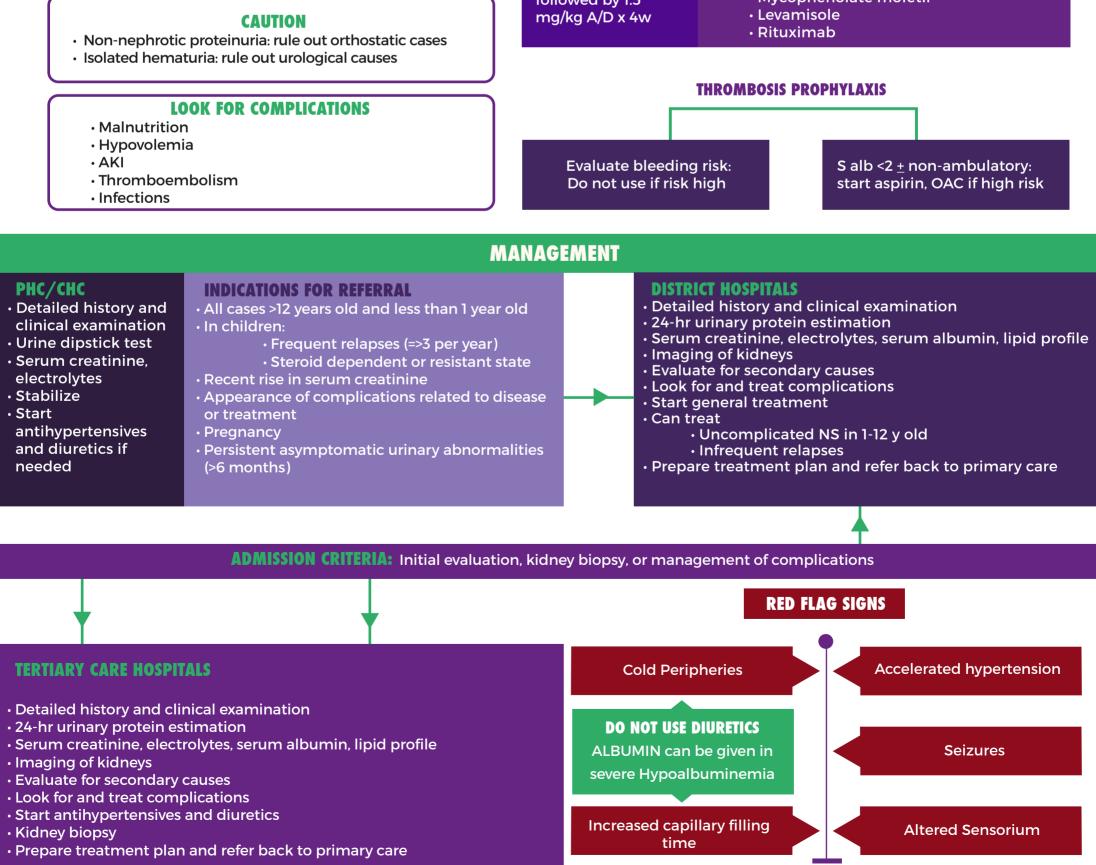
- · Look for latent TB (Mantoux test, Chest X-ray) · Start 6 months INH therapy (5mg/kg day) if asymptomatic Mantoux +ve
- · Be on the lookout for common infections (e.g. peritonitis, pneumonia and skin infections)

Prednisolone 2 mg/kg x 6 w followed by 1.5 mg/kg A/D x 6w

 In case of relapse-Prednisolone 2 mg/kg x 2w followed by 1.5

• Treatment Depends on diagnosis (biopsy, serology)

- Therapeutic choices include Corticosteroid (Prednisolone, IV methylprednisolone)
 - CNIs (cyclosporine/tacrolimus)
 - Cyclophosphamide
 - Azathioprine
 - Mycophenolate mofetil



KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



Ministry of Health and Family Welfare, Government of India

Standard Treatment Workflow (STW) for the Management of **URINARY TRACT INFECTIONS** ICD-10-N39.0

			IV-N55.V				
	D	ETERMINE UTI TYPE		MANAGEMENT		PRIMARY/ SE	CONDARY LEVEL
(Charles and Charles and Char		CYSTITIS/ LOWER UTI ia, urgency, frequency	i notor y	ARE and Examination red flag signs and		• History	
WHAT IS UTI? At least 3 Symptoms (dysuria, frequency, urgency, suprapubic pain) OR Dipstick +ve for leucocyte esterase and nitrite if <3 symptoms	• Fever • Chills • Loin p • Renal • Toxic a • Toxic a	 Chills and rigors Loin pain, pelvic pain Renal angle tenderness Toxic and sick appearance COMPLICATED UTI History of stones, congenital anomalies, obstruction 		 refer Refer special groups Treatment in primary care Acute cystitis females Acute cystitis males Treatment Nitrofurantoin Trimethoprim sulphamethoxazole 		• Examinatio • Tempera	ent UTI es ital nations osuppression on ature, renal enderness,
 OR CUE >10 WBC/HPF in uncentrifuged urine + Urine culture >105 CFU of single species/ml in Mid stream urine (multiple species indicate contamination) WHEN TO DO URINE CULTURE? Complicated UTI Pyelonephritis Special situations All males (except simple cystitis) Children Pregnancy Recurrent UTI (> 2 episodes/ 6 months) 		Rx Men for Nitrofuran TMP/SMX Ciprofloxa Levofloxa Acute cyst All OTHER U Pyelonepl	MALE UTI Acute cystitis/ simple UTI • Rx Men for 7 days • Nitrofurantoin* 100 mg PO BD x 7d • TMP/SMX 1 DS tab PO BD x 7d • Ciprofloxacin 500 mg BD x 7 days • Levofloxacin 750 mg OD x 5 day • Acute cystitis is not to be referred ALL OTHER UTI IN MALES-REFER • Pyelonephritis or complicated UTI • Pelvic/perineal pain (prostatitis)		 Empirical 	ntoin* 100 d 1 DS tab d onse refer	
	· C	atheter associated UTI		*Avoid if GFR <4	5,cau	tion in elderly	
		SYMPTOM	ATIC TREATMEN	1			
	er recommended eg citi atient on nitrofurantoin	rate Phenazopyrid 200mg tid for 2		strogen creams for recu n post menopausal won		Paracetam for pain	ol Cranberry can be used
		RED F Special situations (Child nales except simple cys		• Non respo		within 3 days o	fАВ
<u>ــــــــــــــــــــــــــــــــــــ</u>							
		TERI	IARY LEVEL				
 Send for culture Imaging if no response t Urology services if obstru 		Rx Pyelonephritis/ complicated UTI	Rx pregnancy UTI	Rx all male UTI including prostatitis	r	Rx ecurrent UTI	Rx non-resolving UTI

Empiric Outpatient:

 Urine culture at 1st antenatal visit

Rx of asymptomatic

CATHETER UTI

UTI symptoms+ pelvic

MALES WITH PROSTATITIS

Uncomplicated

• Urine c/s

 Consider initial dose of a parenteral agent

PYELONEPHRITIS

- Ceftriaxone 1-2 g IV/IM x 1
- Gentamicin 5 mg/kg IV/IM x 1 Followed by
- Ciprofloxacin 500 mg PO BD x 7d
- Levofloxacin 750 mg PO OD x 5 d
- Cefuroxime 500 mg PO BD x10-14d
- Amoxy clav x10-14 days
- TMP-SMX 1 DS BD x 7-10 days **Empiric Inpatient :**
- Ceftriaxone 1-2 g IV once daily+ /-AMP
- Gentamicin +/-AMP
- Others as per c/s- Carbapenem, **Piperacillin Tazo**

IV therapy required until afebrile x 48 hrs, then switch to PO If no response in 3 days imaging

bacteriuria/acute cystitis: - Nitrofurantoin 100 mg PO BD x 5-7 d (avoid near-

term) - Cephalexin 500 mg PO QID x 5-7 d

• For asymptomatic

- TMP/SMX 1 DS tab PO BD x 5-7 d (avoid in 1st trimester & near term; supplement with multivitamin containing folic acid)

PREGNANCY UTI

- Check repeat urine c/s 7days after Rx to confirm clearing Repeat urine culture in each antenatal visit
- If recurrent- Antibiotic prophylaxis till term

CAUTI NOT

- recommended • Urinary catheters should be removed as
- soon as not required If indwelling catheter for >2 weeks and is still indicated, replacing the catheter is
- recommended Symptomatic CAUTI - (Fever, back pain, new
- onset delirium, rigors) - Send culture
- Rx as complicated UTI • No role of routine
- antibiotic prophylaxis for prevention

CHILDREN

- pain/ fever Refer
- Urine culture & MSU
- · Digital rectal examtender prostate
- Older >35 yrs-
- Septran DS BD - Levofloxacin 500mg OD, ciproflox 500 mg BD
- Avoid nitrofurantoin Young males-
- Doxy 100mg bd /azithro 1 gm / oflox 300mg BD for chlamydia + Single dose of Ceftrioxone 250mg IM for gonorrhoea • Rx- 6 weeks
- Imaging to rule out abscess

RUII

RECURRENT UTI

- post coital voiding and post coital antibiotic
- Low dose nitrofurantoin 50 mgX 6 months
- Single strength septran x 6 months
- Or norflox 200mg, ciproflox200mg, cephalexin 250mg
- Vaginal cream in post menopausal
- Complicated RUTI
- Urology referral
- Cystoscopy, urodynamics (post menopausal)

ASYMPTOMATIC BACTERIURIA

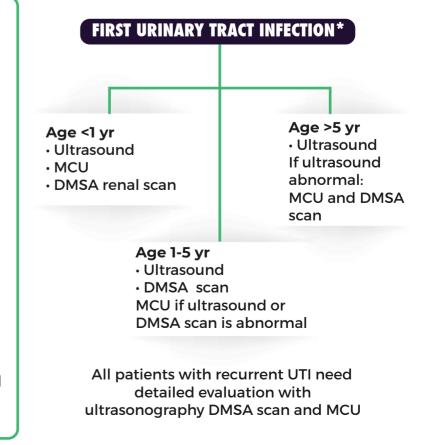
- No symptoms
- Bacteria in urine culture >105CFU/ml
- No treatment required
- Exceptions when you should treat
- Pregnancy
- Before any urological intervention

* Pregnancy UTI, Catheter UTI may also be managed at secondary level.

LONG TERM CONSEQUENCES

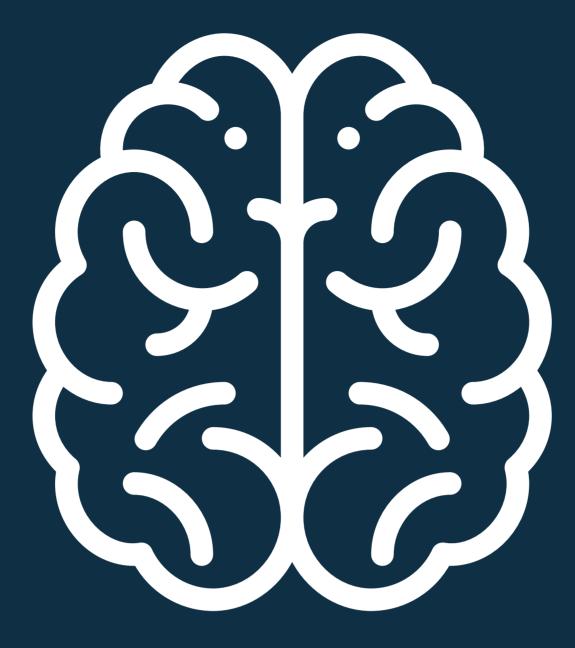
- Renal scars
- Hypertension
- CKD
- Poor quality of life

- Neonates and Infants < lyr - Fever, vomiting, diarrhoea, jaundice, Poor stream TREATMENT
- gentamicin, amikacin, ceftriax one
- ciproflox, amoxyclav
- Upper UTI- 10-14 days
- Lower UTI 7-10 days
- Adolescents 3-5 days
- Upper UTI(PN), infants UTI, recurrent UTI PREVENTION
- Avoid constipation, clean washrooms



🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

- **SYMPTOMS**
- Older children same as adults
- Infants <3months as upper UTI (PN) with IV antibiotics
- Urinary bladder catheterisation for infants with upper tract UTI
- Older children
- Upper UTI- IV antibiotics
- · Lower UTI- oral cefixime, oflox,
- Duration of Rx
- REFER



NEUROLOGY

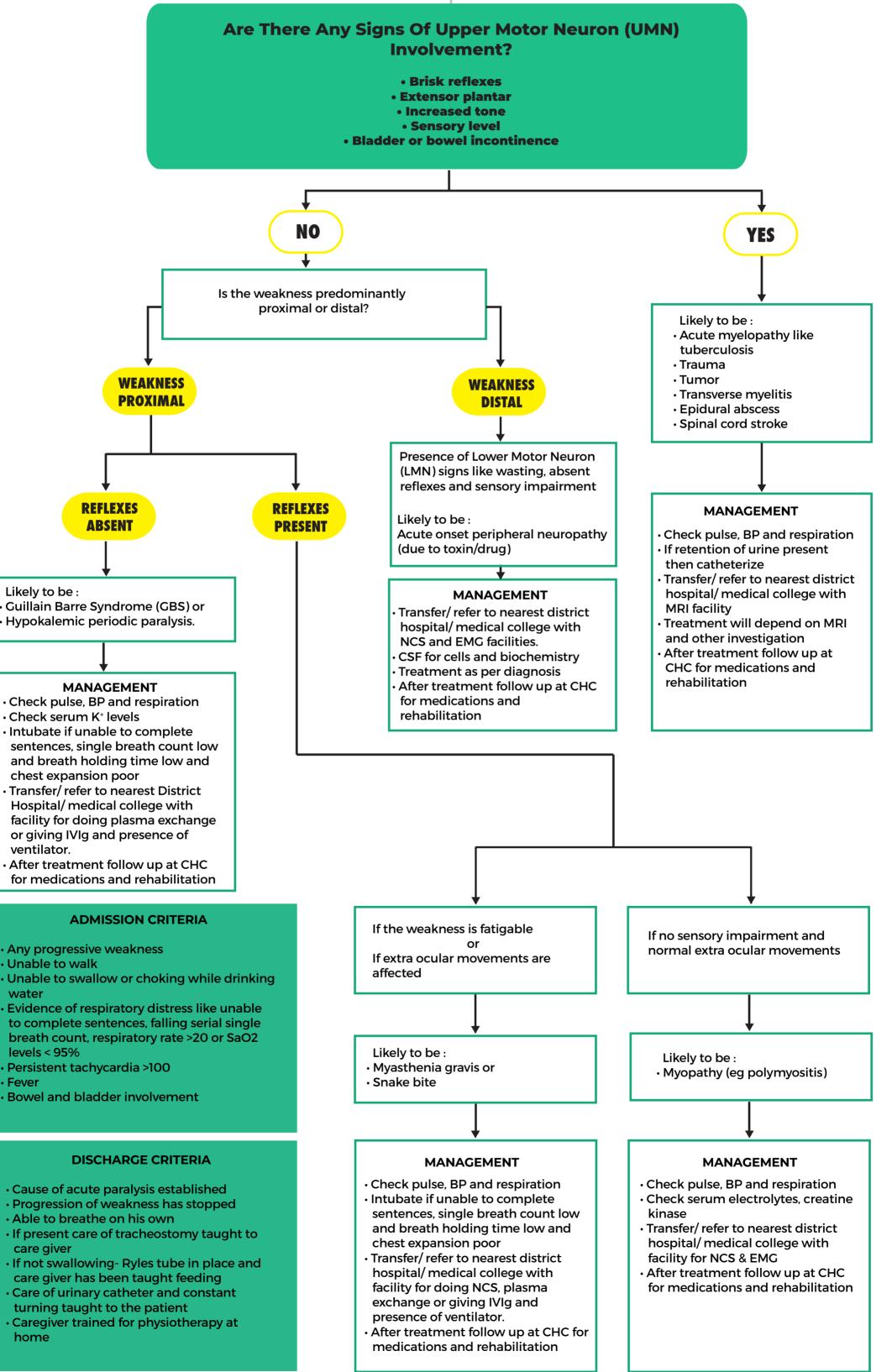


Ministry of Health and Family Welfare, Government of India



Standard Treatment Workflow (STW) for the APPROACH TO ACUTE PARALYSIS ICD 10 G82, G83

PRESENTATION WITH ACUTE ONSET (WITHIN HOURS TO DAYS) PARAPLEGIA OR QUADRIPLEGIA



- Check pulse, BP and respiration
- Check serum K⁺ levels
- Intubate if unable to complete sentences, single breath count low and breath holding time low and chest expansion poor
- Hospital/medical college with facility for doing plasma exchange or giving IVIg and presence of ventilator.
- for medications and rehabilitation

- Any progressive weakness
- Unable to walk
- · Unable to swallow or choking while drinking water
- Evidence of respiratory distress like unable to complete sentences, falling serial single breath count, respiratory rate >20 or SaO2 **levels < 95%**
- Persistent tachycardia >100
- Fever
- Bowel and bladder involvement

- Cause of acute paralysis established
- Able to breathe on his own
- If present care of tracheostomy taught to care giver
- · If not swallowing- Ryles tube in place and
- Care of urinary catheter and constant turning taught to the patient
- Caregiver trained for physiotherapy at

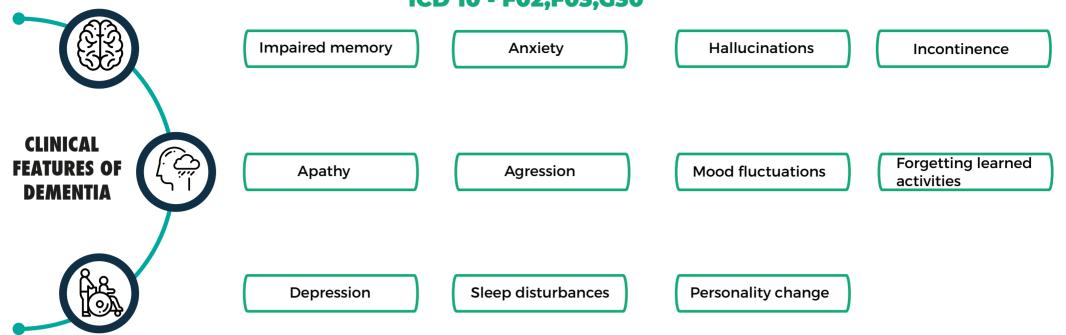
KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES





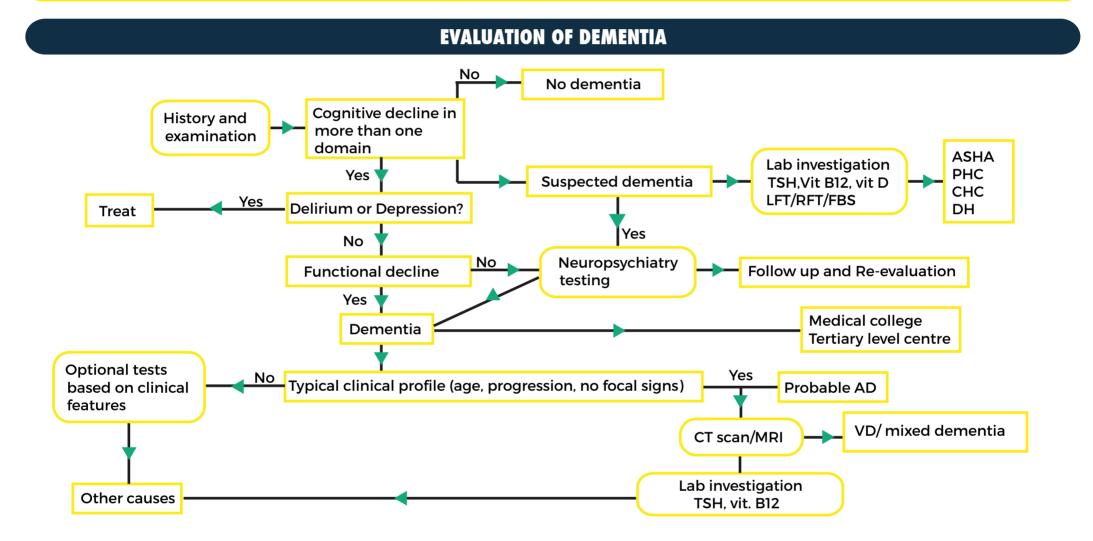
Standard Treatment Workflow (STW) for the Management of

DEMENTIA ICD 10 - F02,F03,G30



IMPORTANT POINTS TO CONSIDER

- Dementia is a complex and variable condition
- No single test will definitively diagnose dementia
- The clinical features if present, should be a change from baseline normal functioning in a middle aged to old person
- Assessment should aim at gathering information about changed behaviours, functional capacity, psychosocial support and medical comorbidities
- History should be taken from a close caregiver, staying with the patient for a longer duration than the appearance of symptoms



FOLLOW UP OF DIAGNOSED & TREATED PATIENTS INTERVENTION MATRIX FOR DEMENTIA ACROSS PLATFORMS OF CARE

PRIMARY HEALTH CENTRE (MEDICAL OFFICER)

- Diagnose dementia after detailed history
- Screening for:
- Treatable causes of dementia thyroid disorders, B-12 deficiency, subdural hemorrhage.
- Depression.
- Vascular risk factors
- · Lab investigations- CBC, biochemistry, liver function tests, hemogram, lipid profile, TFT, VDRL, vit B12 level, vit **D** level
- Referrals for MRI/CT
- Initiation of treatment/drugs; treatment for co-morbid conditions (including depression, vision, hearing deficits and gait problems), thyroid, arthritis.
- Initiate therapy for vascular risk factors
- Encourage healthy lifestyle
- Assess for palliative care
- · Learn and share facts about dementia to provide immediate need to the person with severe dementia
- Follow up and monitor for side effects of drugs/ red flags in patient/ signs of danger
- · Follow-up of difficult patients under the guidance of higher centre.

DISTRICT HOSPITAL (SPECIALIST- PHYSICIAN/ **GERIATRIC SPECIALIST / NEUROLOGIST / PSYCHIATRIST**)

- · Careful evaluation of all the referral patients of dementia
- Screening for treatable causes for dementia including normal pressure hydrocephalus, B12 deficiency, hypothyroidism, chronic meningitis
- Neuroimaging CT/MRI- to rule out subdural hematoma/ tumors/ NPH(surgically remediable causes of rapid cognitive decline)
- Lab investigations- CBC, liver function tests, biochemistry, hemogram, lipd profile, vit D levels, TFT, VDRL, retrovirus after counselling (whenever feasible and high index of suspicion)
- All the points mentioned in PHC to be followed if patient presents to a DH
- Upward referral linkages with tertiary care and downward referral with PHC.
- Encouraging patient and caregiver participation in an ongoing support program for them.
- Avoid antipsychotics until necessary
- Interaction with, training of MOs at PHC/UPHC and ongoing clinical support and supervision

REASONS FOR REFERRAL

- Not responding to adequate dose and duration of prescribed medications
- Presence of red flags

RED FLAGS

- **Fever**
- **Rapid progression**
- Seizures
- **Recent head**
- injury Alcoholism and falls

MEDICATIONS RECOMMENDED FOR USE FOR ALZHIEMERS DEMENTIA

FOR COGNITION

- Donepezil: 5 mg once after breakfast x 1 month, then 10 mg after breakfast to continue If any side effect/ not tolerating: **Rivastigmine** to be used start dose 1.5 mg BD / 1 month then 3 mg BD x 1 month, then 4.5 mg BD x 1 month, then 6 mg twice after meals only x 1 month.
- Memantine: in moderate to severe dementia 5 mg BD x 1 month, then 10 mg BD to continue.
- Galantamine: 8 mg BD if not tolerating 1

🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

This STW has been prepared by national experts of India with feasibility considerations for various levels of healthcare system in the country. These broad guidelines are advisory, and are based on expert opinions and available scientific evidence. There may be variations in the management of an individual patient based on his/her specific condition, as decided by the treating physician. There will be no indemnity for direct or indirect consequences. Kindly visit our web portal (stw.icmr.org.in) for more information. © Indian Council of Medical Research and Department of Health Research, Ministry of Health & Family Welfare, Government of India.

FOR DEPRESSION

FOR AGITATION

- Identification of triggers
- Non pharmacological interventions

Escitalopram 10 mg





Standard Treatment Workflow (STW) for the Management of EPILEPSY ICD 10 - G40

CLINICAL	Episodic, few secs-mins	Abnormal jerky movements	Loss of consciousness/ awareness	Blank staring	Episodes could be single with high risk of recurrence. Prolonged motor convulsion of > 5
FEATURES	Sudden drops/ falls, brief jerks	With or without urine/ stool incontinence, tongue bite, drooling	Bizzare activity out of context	Any other episode lasting for few mins (usually<5 mins)	mins with loss of consciousness - STATUS EPILEPTICUS (SE) - MEDICAL EMERGENCY

PRIMARY HEALTH CENTRE (MEDICAL OFFICER)

- Clinical diagnosis of epilepsy: detailed history from an eyewitness
- Differentiate between provoked seizures and epilepsy (provoked due to fever, acute CNS insult, antibiotics, and metabolic causes)
- Laboratory investigations: CBC, liver function tests, routine biochemistry, hemogram, lipid profile, vit D levels, TFT (whichever feasible)

Initiation of treatment:

- Treat the patient if patient has epilepsy (2 or more episodes of unprovoked seizures)
- Treat a single seizure if risk of recurrence is high as in patients with focal seizures, mentally retarded, neurological deficits having family history of seizures abnormalEEG, neuroimaging
- Anti Epileptic Drug (AED broad spectrum, low dose, start low go slow, except status epilepticus)
- Emergency medical care of status epilepticus
- Treatment counselling: side effects/toxicities of drugs, red flags, importance of adherence, maintaining treatment diary
- Advice on prevention of seizures: regular medication, sleep 7-8 hrs, avoid excess TV/mobile/ photic stimulation, regular diet, lifestyle choices(avoid alcohol)
- Evaluate any possibility of superimposed non-epileptic seizure
- Training of MLP/ANM/ASHA on epilepsy
- For excessive alcohol use, refer to ANM/MLP where psychosocial interventions are carried out for substance use disorders
- Follow up visits for treatment monitoring & difficult patients under neurologist at STC centre
- Basic management of co-morbidities (behaviour, cognition, reproductive health, bone health)
- Alert to signs of abuse and neglect
- Maintain upward referrals with paediatrician/physician at DH

AED- BROAD SPECTRUM (GENERALIZED SEIZURES)

Sodium Valproate (avoid in women of child bearing age unless non responsive to other drugs)

DOSE (MAINTENANCE: MG/D)

Starting dose :200mg TDS Maintenance Dose: 600-2400

Starting dose: 25mg HS (Lower dose with VPA) Maintenance Dose: 100-300

REASONS FOR REFERRAL

(centres with specialists like paediatrician, neurologist)

- Redflag Signs
- Progressive problems, rapid
 appearance of new symptoms
- Recent injury
- Symptoms appearing after alcohol binge
- Status epilepticus after stabilization
- Non response to adequate dose and duration of medication
- Serious side effects
- Neuroimaging

RED FLAG SIGNS

- Fever Headache
- Vomiting
- Altered Sensorium
- Severe Giddiness
- Loss of function of body

DISTRICT HOSPITALS

- Careful evaluation of all referral patients, provide specialized management for patients and refer back to PHC for follow up of management
- Maintain communication, ongoing clinical support and supervision of MOs at PHC
- Laboratory investigation - CBC. liver function
- tests, antiepileptic drug levels, routine biochemistry, hemogram, lipid profile,
- vit D levels, TFT, CT brain (when necesary)
- Monitor side effects of AED
- Clinical Psychologist: counselling health services for persons with epilepsy or upon referral from PHC/UPHC

ADVERSE EFFECTS

Anorexia, wt gain, nausea, vomiting, tremors, hair loss, PCOS, thrombocytopenia

Sedation, ataxia, dizziness, skin rash, SJS (lower risk with slow titration)

Lamotrigine

ge	Maintenance Dose: 100-300	risk with slow titration)		
Levetiracetam	Starting dose: 250mg BD Maintenance Dose: 1000-3000	Somnolence, dizziness, cognitive slowing, psychosis		
Topiramate	Topiramate Starting dose: 25mg OD Maintenance Dose: 100-400			
AED (focal seizures)				
Carbamazepine	Starting dose: 100mg BD Maintenance dose: 400-1200	Sedation, dizziness, ataxia, skin rash, SJS, hyponatremia, seizure worsening in some situations		
Oxcarbazepine	Starting dose: 150mg BD Maintenance dose: 600 to 1800	Sedation, dizziness, ataxia, headache, hyponateremia, skin rash		
Phenobarbitone Can be used for generalized also	Starting dose: 30mg HS Maintenance dose: 60-180	Sedation, ataxia, depression, memory problems, hyperactivity in children, skin rash		
Phenytoin	Starting dose: 200mg HS Maintenance dose:200-400	Ataxia, sedation, gum hyperplasia, coarsening of facial features, hirsutism, memory problems, osteomalacia & bone loss, skin rash		
Folic Acid 5 mg/day to be added along	with AEDs in all women of child bearing age. Polytherapy and	d valproate to be avoided in women with epilepsy		
IMPENDING SE	ESTABLISHED SE	REFRACTORY SE		
5 MIN FIRST ABCS TO	30 MIN BE DONE FROM WHEN YOU SEE PATIENT SIMULTANEOU	60 MIN 2 IV drugs fail (Benzo + IV AED) JSLY WITH MEDICATION		
FIRST ABCS TO		JSLY WITH MEDICATION		
FIRST ABCS TO	BE DONE FROM WHEN YOU SEE PATIENT SIMULTANEOU	ICU IV Midazolam loading 0.2 mg/kg OR CIV 0.05-0.5 mg/kg/hr		
FIRST ABCS TO Out of Hospital/home : Buccal/Intranasal	BE DONE FROM WHEN YOU SEE PATIENT SIMULTANEOU IMDZ with acute repititive seizures/status (0.3-0.5 mg/kg)	ICU IV Midazolam loading 0.2 mg/kg OR CIV 0.05-0.5 mg/kg/hr (can go up to 2 mg/kg/hr) Taper gradually after seizure stops		
FIRST ABCS TO Out of Hospital/home : Buccal/Intranasal EMERGENCY ROOM IV Lorazepam up to 0.1 mg/kg @ 2mg/m OR IV Midazolam 0.1-0.2 mg/kg bolus or 0.0	BE DONE FROM WHEN YOU SEE PATIENT SIMULTANEOU IMDZ with acute repititive seizures/status (0.3-0.5 mg/kg) in	ICU IV Midazolam loading 0.2 mg/kg OR CIV 0.05-0.5 mg/kg/hr (can go up to 2 mg/kg/hr) Taper gradually after seizure stops (preferably as evidenced by EEG)		
FIRST ABCS TO Out of Hospital/home : Buccal/Intranasal EMERGENCY ROOM IV Lorazepam up to 0.1 mg/kg @ 2mg/m OR	BE DONE FROM WHEN YOU SEE PATIENT SIMULTANEOU IMDZ with acute repititive seizures/status (0.3-0.5 mg/kg) in 5-0.5 mg/kg/hr in CIV	ICU IV Midazolam loading 0.2 mg/kg OR CIV 0.05-0.5 mg/kg/hr (can go up to 2 mg/kg/hr) Taper gradually after seizure stops		
FIRST ABCS TO Out of Hospital/home : Buccal/Intranasal EMERGENCY ROOM IV Lorazepam up to 0.1 mg/kg @ 2mg/m OR IV Midazolam 0.1-0.2 mg/kg bolus or 0.0 OR IV Diazepam upto 0.25-0.4 mg/kg over 2 . Phenytoin @50 mg/min 20 mg/kg plus 10 mg/kg if seizures do not sto . If seizures not controlled or contra Intravenous Valproate 25-40 mg/kg . If CI to above two; Phenobarbitone prepared to Intubate and ventilate	BE DONE FROM WHEN YOU SEE PATIENT SIMULTANEOU IMDZ with acute repititive seizures/status (0.3-0.5 mg/kg) in 5-0.5 mg/kg/hr in CIV -3 min repeat op in 15-20 min indiction (CI) to PHT g @3-6 mg/kg/min 20 mg/kg IV @ less than 5-60 mg/min but be	JSLY WITH MEDICATION ICU IV Midazolam loading 0.2 mg/kg OR CIV 0.05-0.5 mg/kg/hr (can go up to 2 mg/kg/hr) Taper gradually after seizure stops (preferably as evidenced by EEG) Thiopental 5-7 mg/kg IV bolus further 50 mg until seizures controlled		
FIRST ABCS TO Out of Hospital/home : Buccal/Intranasal EMERGENCY ROOM IV Lorazepam up to 0.1 mg/kg @ 2mg/m OR IV Midazolam 0.1-0.2 mg/kg bolus or 0.0 OR IV Diazepam upto 0.25-0.4 mg/kg over 2 . Phenytoin @50 mg/min 20 mg/kg plus 10 mg/kg if seizures do not sto . If seizures not controlled or contra Intravenous Valproate 25-40 mg/kg . If CI to above two: Phenobarbitone prepared to Intubate and ventilate	BE DONE FROM WHEN YOU SEE PATIENT SIMULTANEOU IMDZ with acute repititive seizures/status (0.3-0.5 mg/kg) in 5-0.5 mg/kg/hr in CIV -3 min repeat op in 15-20 min indiction (CI) to PHT g @3-6 mg/kg/min 20 mg/kg IV @ less than 5-60 mg/min but be	JSLY WITH MEDICATION ICU IV Midazolam loading 0.2 mg/kg OR CIV 0.05-0.5 mg/kg/hr (can go up to 2 mg/kg/hr) Taper gradually after seizure stops (preferably as evidenced by EEG) Thiopental 5-7 mg/kg IV bolus further 50 mg until seizures controlled 3-5 mg/kg/hr for only 48 hours OR Propofol IV loading 2-5 mg/kg CIV 1-15 MG/KG/HR OR Pentobarbital IV upto 10 mg/kg @ <0.2-0.4 mg/kg/min CIV 0.5-2 mg/kg/h OR Ketamine bolus 1.5 mg/kg CIV 0.01-0.05 mg/kg/h max 10mg/kg/hr * to be		

Airway, blood pressure, temperature, intravenous access, electrocardiography, CBC, glucose, electrolytes, AED levels, ABG, oximetry, tox screen, central line If alcoholic- thiamine & glucose, if diabetic GLUCOTEST/blood sugar & glucose IV. MUST INFORM CONSULTANT ON CALL

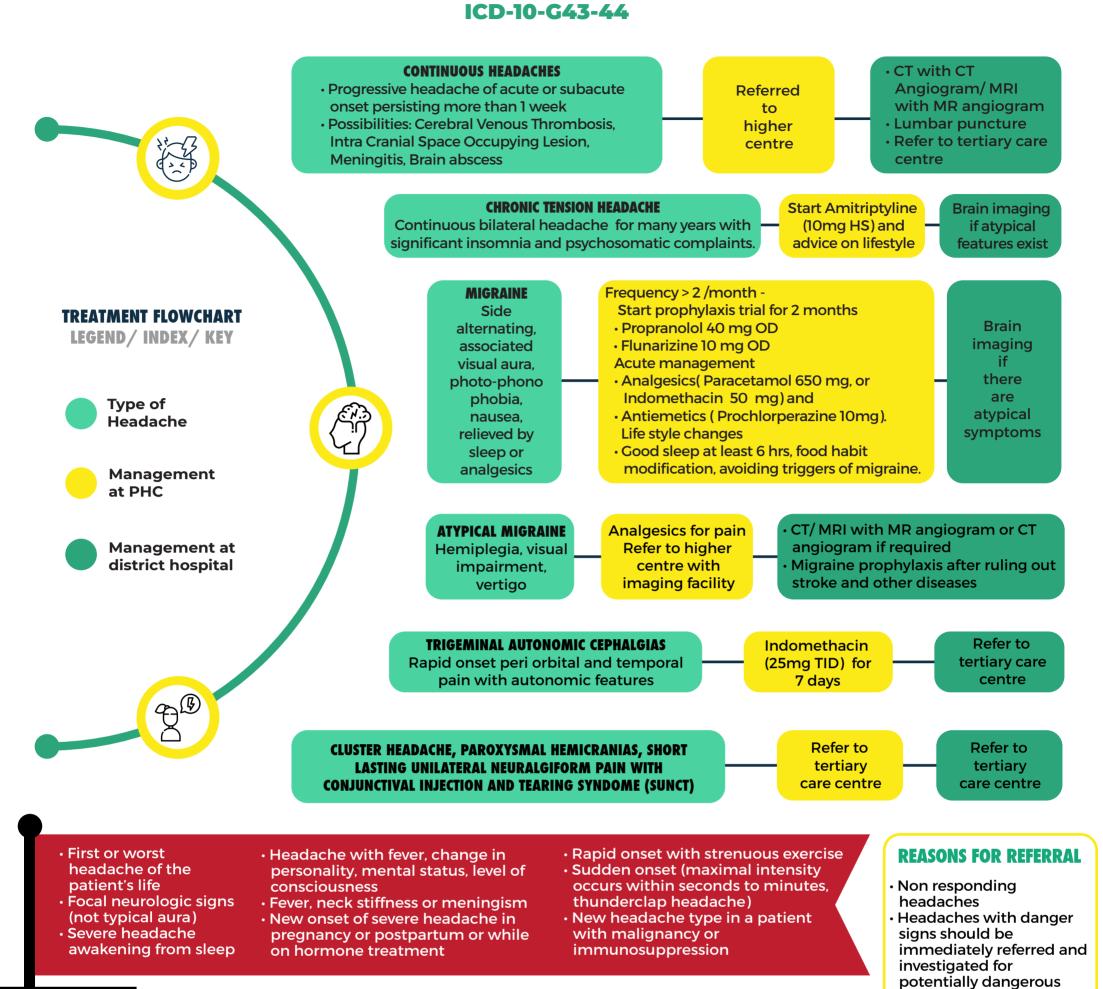
KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

सत्यमेव जपते

Department of Health Research Ministry of Health and Family Welfare, Government of India



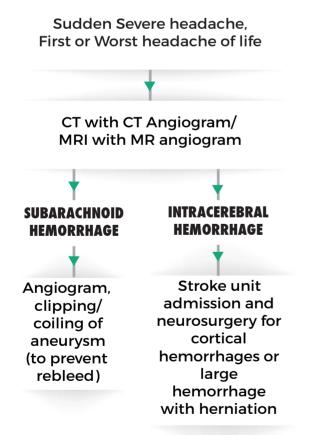
Standard Treatment Workflow (STW) for the Management of **HEADACHE**

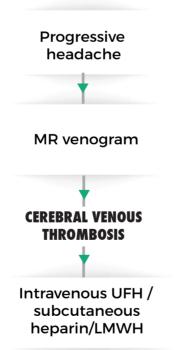


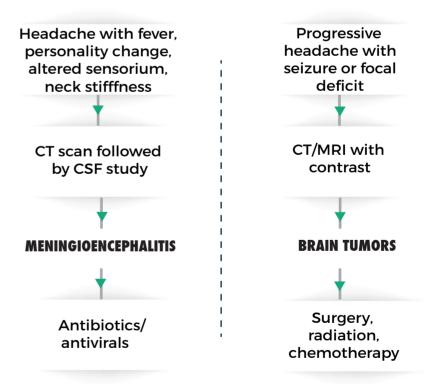


conditions

TREATMENT OF MAJOR CATASTROPHIC HEADACHES AT TERTIARY CENTRE







INDICATIONS FOR ADMISSION

- Patient with unrelenting headache
- Immunosuppressed patients
 with continuous headache,
- First ever headache with worsening intensity,
- Progressive headache with other systemic disease
- Severe symptomatic primary headache disorders

CRITERIA FOR DISCHARGE

- Primary headache disorderssymptomatically improved severe episode of headache due to primary headache disorder can be discharged
- Secondary headache disorderssecondary headache disorders with essential work up, diagnosis and treatment as per individual case can be discharged

FOLLOW UP OF HEADACHE PATIENTS

CAUSES OF HEADACHE	TREATMENT OF HEADACHE
Intra cerebral hemorrhage	Good control of blood pressure
Seizures	Antiepileptic medications
Cerebral venous sinus thrombosis	Follow up of anticoagulation
Migraine	Give prophylaxis for adequate duration of time and taper after remission

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

REFERENCES

1. Hainer BL, Matheson EM. Approach to acute headache in adults. American family physician. 2013 May 15;87(10).

2. https://www.uptodate.com/contents/evaluation-of-headache-in-adults

ABBREVIATIONS

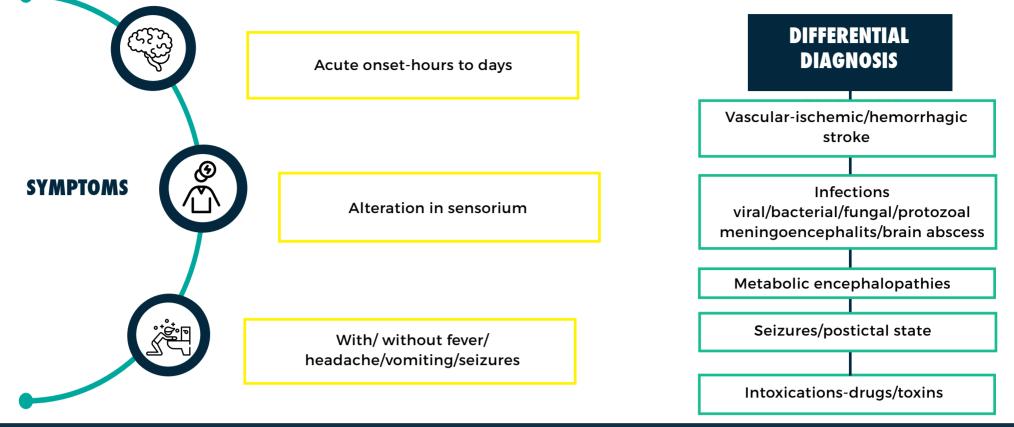
CSF: Cerobrospinal Fluid, UFH: Unfractionated Heparin, LMWH: Low Molecular Weight Heparin





Standard Treatment Workflow (STW) for the Management of **NEUROINFECTIONS**

ICD-10-G03.9



AT PRIMARY CARE LEVEL

ESSENTIAL

Check Airway/Breathing/Circulation

Rule out circulatory shock, ongoing convulsions and hyperthermia/hyperpyrexia(core temperature > 40.5°C or hypothermia(< 36.5°C)

Establish IV access-urgent blood for hemogram/sugar/electrolytes/malaria testing-peripheral smear/rapid antigen detection

Correct hypoglycemia (blood sugar 50 mg/dl) with IV 100ml of 25% dextrose solution

If seizing- IV/IM Lorazepam 0.1 mg/kg followed by loading with Phenytoin 20 mg/kg weight at a rate of 50 mg/minute

When IV access not available-intra nasal or buccal Midazolam 0.2 mg/kg /intra rectal Diazepam 0.3-0.4 mg/kg

Urgent referral to higher centres with intensive care facilities

AT SECONDARY CARE LEVEL(TALUK, DISTRICT) HEADQUARTERS HOSPITAL

NOT RECOMMENDED

- Stomach wash
- Inj Mannitol
- Inj Steroids

CRITERIA FOR REFERRAL

Altered sensorium/seizures /focal deficits/hemodyna mic instability -where imaging and ICU management are required.

In addition to all the steps given above :

teps given above :

Establish and maintain airway: Intubate if GCS<8, impaired airway reflexes, abnormal respiratory pattern, signs of raised ICP, oxygen saturation <92% despite high flow oxygen, and fluid refractory shock

Inj Thiamine 100 mg IV

Stomach wash/activated charcoal administration-if history or suspicion of drug overdose/ non corrosive poison intake

Start treatment for cerebral malaria-first dose of IV Artesunate 2.4 mg/kg OR Quinine 20 mg/kg bolus

Emergency CT/referral to centre with 24 hour CT facilities

CRITERIA FOR REFERRAL

Altered sensorium/seizures/focal deficits/hemodynamic instability -where imaging and ICU management are required.

 \cdot If no definite diagnosis achieved after preliminary investigations

AT TERTIARY CARE HOSPITALS-SELECTED DISTRICT HOSPITALS/MEDICAL COLLEGES

- Neuroimaging-MRI/CT with contrast to rule out abscess/herniations.
 If abscess-emergency neurosurgical consultation for favour of aspiration –open/stereotactic
- Blood cultures-aerobic/anaerobic

• CSF analysis-biocehmistry/cytology/gram staining/culture-bacterial , AFB and fungal/viral PCR/TB-PCR/fungal antigen

 Empirical antibiotic (within 30 minutes of arrival) If suspecting pyogenic meningitis-Inj Ceftriaxone 2 g+ Inj Vancomycin 500 mg+ Inj Ampicillin 2 g if older than 50 years or immunocompromised+ Inj Dexamethasone 8 mg IV Continue empirical treatment till culture yields causative organism,then tailor treatment as per sensitivity reports for 10-14 days. Steroids to be stopped after 48 hours,unless any other compelling indications-adrenal insufficiency/TBM 		Viral-Herpes simplex/Zoster • Inj Acyclovir 500 mg IV 8 hourly for 10 days	Cerebral malaria Inj Artesunate 2.4 mg/kg IM or IV 3 doses 12 hours apart and then OD / Inj Quinine 20 mg/kg IV stat followed by 10mg/kg TDS till patient can take orally,then oral Artesunate+Pyrimethamine /Sulphadoxine fo 3 days OR oral Quinine 10 mg/kg TDS for total 7 days + Doxycycline 3 mg/kg OD for 7 days.			
COMPLICATIONS						
Raised ICP	SIADH		Vasculitis		Hydrocephalus	
*If uncomplicated-back referral to Secondary care centre for completing treatment regimen/monitoring.						
CRITERIA FOR DISCHARGE						
Afebrile,hemodynamically stable,seizure Diagnosis and treatmer free >48 hours initiate		•	Continuation of treatment with monitoring can be ensured for the prescribed duration.			
KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES						

This STW has been prepared by national experts of India with feasibility considerations for various levels of healthcare system in the country. These broad guidelines are advisory, and are based on expert opinions and available scientific evidence. There may be variations in the management of an individual patient based on his/her specific condition, as decided by the treating physician. There will be no indemnity for direct or indirect consequences. Kindly visit our web portal (**stw.icmr.org.in**) for more information. **©** Indian Council of Medical Research and Department of Health Research, Ministry of Health & Family Welfare, Government of India.

DESIRABLE

Neuroimaging-CT with contrast -to rule out hemorrhage/infarcts/focal edema or lesions Blood cultures aerobic/anaerobic

First dose of empirical treatment of pyogenic meningitis-Inj Ceftriaxone 2 g + Inj Vancomycin 500 mg.

Add Inj Ampicillin 2 g if older than 50 years / immunocompromised along with Inj Dexamethasone 8 mg Fundus examination,CSF study to rule out

meningoencephalitis-if imaging rules out any mass lesions/herniations.

Urgent referral to higher centres with Intensive care facilities



Department of Health Research Ministry of Health and Family Welfare, Government of India



Standard Treatment Workflow (STW) for the Management of **STROKE**

ICD-10-163, 164



WHAT IS STROKE?

An episode of neurological dysfunction caused by focal cerebral, spinal, or retinal infarction or haemorrhage



Numbness or weakness, especially on one side of the body

SYMPTOMS

- Loss of consciousness or altered consciousness
- Decreased vision in one or both eyes
- Language difficulties, either in speaking or understanding
- Difficulty walking; loss of balance or coordination
- Confusion or loss of memory
- Swallowing difficulties
- Paralysis of any part of the body, including face
- Sudden, severe headache with no known cause
 Neck pain
- Nausea and vomiting

WARNING SIGNS (BEFAST)

- **B**ALANCE : Loss of balance or coordination
- EYES : Sudden blurred or double vision/ sudden, persistent vision trouble
- FACE : Deviation at the angle of the mouth
- ARM : Arm Drift
- **S**PEECH : Slurred speech or the inability to speak or understand
- TIME : Act fast
- **S**udden new onset of headache or loss of consciousness
- **S**udden giddiness, vomiting and imbalance

	TYPES OF STROKE					
Ischemic stroke Focal cerebral, spinal, or retinal infarction	Intracerebral haemorrhage Focal collection of blood within the brain parenchyma or ventricular system that is not caused by trauma	Subarachnoid haemorrhage Bleeding into the subarachnoid space a cerebral venous structure		nrombosis of I venous	Transient Ischemic Attack (TIA) Transient episode of neurologic dysfunction caused by focal cerebral, spinal cord, or retinal ischemia, without acute infarction	
PRELIMINARY MANAGEMENT INVESTIGATIONS						
 Assess and manage ABCs Initiate cardiac monitoring Maintain O2 saturation >94% Establish IV access Determine blood glucose and treat accordingly Determine time of symptom onset or last known normal, and obtain family contact information, preferably a cell phone Triage and RAPID TRANSFER of patient to nearest district hospital with CT Scan facility or Stroke center with facility for thrombolysis Referal hospital to be notified to handle the referred patient with stroke 		• CT Scan head • CTA		DESIRA • CTA • Echocardi		OPTIONAL • MRI/MRA • Holter monitoring
MANAGEMENT						

STROKE ONSET TIME: <4.5 HOURS

ISCHEMIC: * IV tPA (0-4.5 hrs) or

endovascular treatment according to eligibility and availability

HAEMORRHAGIC:

- · Dysphagia assessment,
- · Blood pressure/blood sugar monitoring and IV fluids.
- Prevention of Pneumonia
- Prophylaxis for deep venous thrombosis etc, monitor and record ECG

* RECOMMENDED DIAGNOSTIC STUDIES			
ALL PATIENTS	SELECTED PATIENTS		
 Noncontrast brain CT or brain MRI Blood glucose Oxygen saturation Serum electrolytes/renal function tests Complete blood count, including platelet count Markers of cardiac ischemia BT, CT, Prothrombin time/INR Activated partial thromboplastin time ECG FLP and carotid doppler (ischemic stroke) 	 TT and/or ECT if it is suspected the patient is taking direct thrombin inhibitors or direct factor Xa inhibitors Liver function tests Toxicology screen Blood alcohol level Pregnancy test Arterial blood gas test (if hypoxia is suspected) Chest radiography (if lung disease is suspected) Lumbar puncture (if subarachnoid hemorrhage is suspected and CT scan is negative for blood) Electroencephalogram (if seizures are suspected) 		

STROKE ONSET TIME: >4.5 HOURS

Rapid Assessment, CODE Stroke, Blood pressure and Blood Sugar monitoring, NIHSS, Intravenous lines Endovascular treatment with Mechanical thrombectomy using stent retriever (4.5 hrs to 24hrs) according to eligibility

SECONDARY PREVENTION

Aspirin (in ischemic stroke) Antihypertensives Antidiabetics Lipid lowering agents

REHABILITATION

Physiotherapy Speech Therapy Occupational Therapy Vocational training

DISCHARGE PLANNING

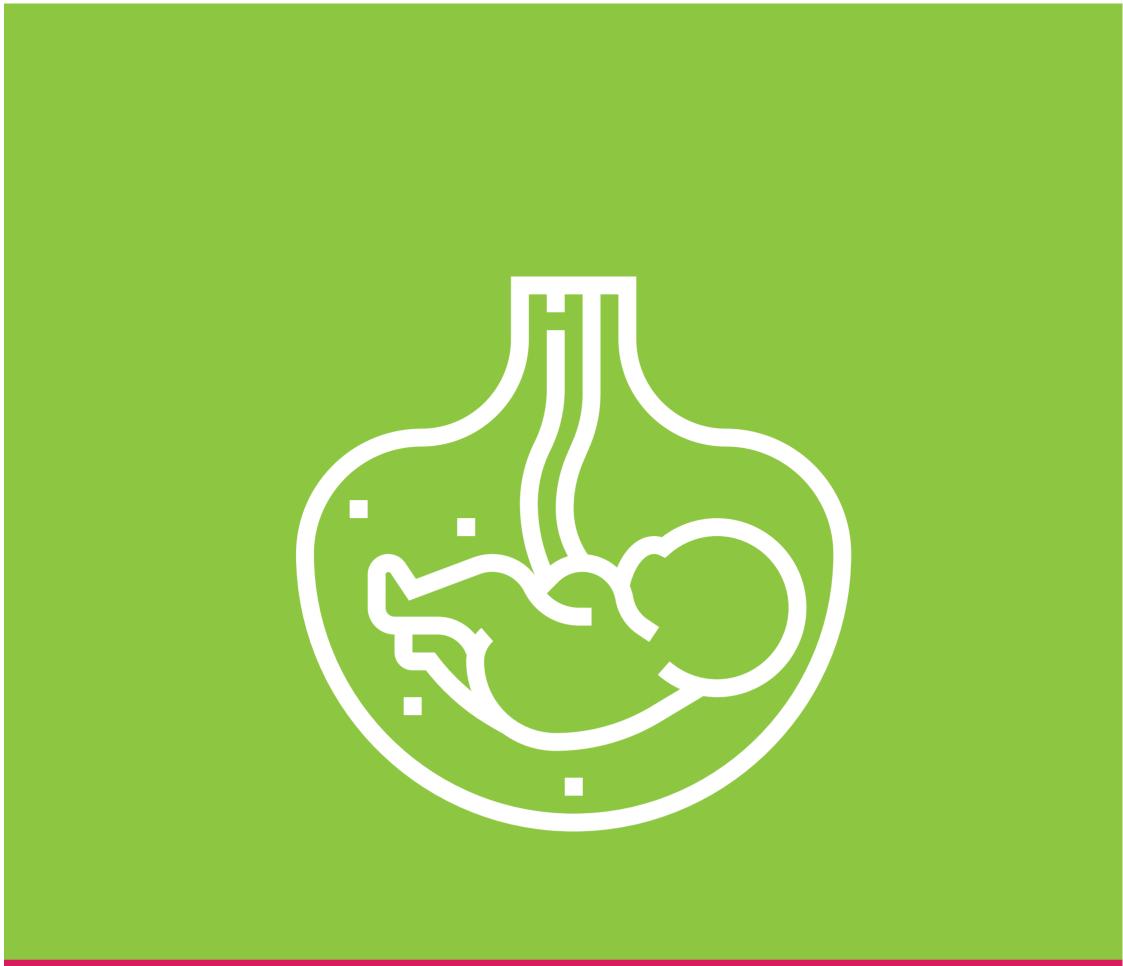
(checklist : drugs, diet, compliance, exercises, health education)

FOLLOW UP at 2nd week, 1st month, 3rd month and 6th month

STROKE UNIT MANAGEMENT

- Medical and Nursing staff : control of blood pressure; control of diabetes; swallow assessment; DVT prophylaxis; antiplatelet drugs
 Rehabilitation staff:
 - » Acute phase: basic bed mobility, transfer techniques, communication training, prevention of complications
 - » Subacute and chronic phase: mobility, gait and balance training, training of activities of daily living (grooming, eating, dressing etc), bowel/bladder training, perceptual and cognitive rehabilitation, provision of assistive devices.

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES









Standard Treatment Workflow (STW) for ANTE-NATAL MANAGEMENT OF NORMAL PREGNANCY

FIRST VISIT (PREFERABLY IN FIRST TRIMESTER)

ASK			EXAMINE	INV	ESTIGATIONS	DO	
 Age LMP Parity & obstetric history Any complaints especially excessive nausea & vomiting/ bleeding PV H/o medical illness : diabetes, hypertension, cardiac problem, epilepsy or any other chronic illness Consanguinity, multiple pregnancy H/o blood transfusion and H/o prior surgical intervention Personal history : tobacco/ alcohol intake Family history : diabetes, hypertension, genetic disorders/ congenital problems, multiple pregnancy, infections including tuberculosis 		 Height, weight Calculate BMI Pallor, Jaundice, Pedal edema Pulse, BP, RR, temperature Thyroid Breast Respiratory and CVS exam - ination P/A examination, P/S and P/V examination # If woman presents with bleeding per vaginum do P/A & P/S to confirm amount of bleeding & rule out local causes. All such cases to be referred to CHC or higher centre 	 ESSENTIAL TESTS Hemoglobin Urine R & M ABO & Rh grouping DESIRABLE TESTS VDRL/ RPR HIV HBsAg WHO OGTT/ DIPSI test for diagnosis of GDM TSH in high risk cases (BOH, goiter, obesity or residing in iodine deficiency prone areas) OPTIONAL TESTS* Aneuploidy screen* by USG & double marker 		 UPT if in doubt Fill up MCH protection card or ANC card, make entry on RCH portal & generate RCH number (in public sector) Give filled MCH protection card & safe motherhood booklet to woman Give Tab Folic Acid daily Give first dose of tetanus toxoid 		
			SECOND VISIT (SECON	ND TRIMESTER)			
ASK • Any com- plaints since last visit • Quickening and/ or fetal movements • Adherence to medications	EXAMINE • Weight • Pallor • Pedal edema • Pulse, BP in sitting position • P/A examination for fundal height	 Hemo Urine DESIRA USG (malfo WHO elapse OPTION Quadr 	albumin BLE TESTS Level II between 18-20 weeks for gros rmations) OGTT/ DIPSI test if >24weeks & at leas ed after 1st test IAL TESTS* ruple test as per availability	INVESTIGATIONS DC TESTS • IFA tablet one (if Hb >11g obin • IFA tablet one (if Hb >11g oumin • Calcium carbonate 500 250 mcg tablet twice data • Calcium Carbonate and together output • Calcium Carbonate and together TT/ DIPSI test if >24weeks & at least 4 weeks have • Single dose of Albendaz after 1st test • Ensure compliance for it treatment			
	THIRD (28 – 34 WEEKS) AND FOURTH VISIT (36 - 40 WEEKS)						

ASK

EXAMINE • Same as above

Same as

- above

Auscultate FHS

 Measurement of abdominal girth and Symphysiofundal

Height

INVESTIGATIONS

- Hemoglobin
- Urine albumin
- Optional USG for fetal
- growth and liquor

• Refer to higher centre if any discrepancy between fundal height and period of

DO

If non compliant or Hb < 9g% give parenteral iron sucrose therapy (not > 200mg

at one time & not > 3 times a week) and refer patient with Hb < 7g% to higher

· Continue IFA and calcium tablets and ensure compliance

DANGER SIGNALS FOR PATIENT TO REPORT TO HEALTH FACILITY

- Fever
- Persistent vomiting
- Abnormal vaginal discharge

HIGH RISK PREGNANCY

- Any H/o medical illness, previous caesarean section, past obstetric mishap or congenital malformation Past H/o PPH

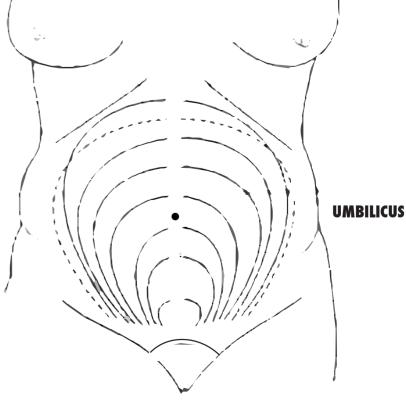
- ry
- fe

centre

gestation

 Palpitations, easy fatigability and breathlessness at rest and/ or on mild exertion. Generalized swelling of the body/ puffiness of the face Vaginal bleeding Decreased or absent fetal movements at > 28 weeks gestation Leaking of watery fluid per vaginum (P/V) Severe headache/ blurring of vision/ convulsion Passing lesser amounts of urine and/ or burning sensation during micturition Itching all over the body 	 Malnouri Hemoglo BP > 140/ APH Discrepa GDM/ ovo Multiple Malprese 	
COUNSELLING AT ALL LEVELS FOR : • Timing and place of next ANC visit based on presence or absence of risk • Rest, nutrition, balanced diet and exercise • Counselling for HIV testing • Danger signs • Institutional delivery • Birth preparedness • Early & exclusive breastfeeding for six months • Post partum contraception	factor	 • •
ASSESSMENT OF FUNDAL HEIGHT & ITS CORRELATION W GESTATIONAL AGE	ITH	

- At 12th week : Just palpable above the symphysis pubis
- At 16th week : At lower one-third of the distance between the symphysis pubis and umbilicus
- At 20th week : At two-thirds of the distance between symphysis pubis and umbilicus
- At 24th week : At the level of umbilicus
- At 28th week : At lower one-third of the distance between the umbilicus and xiphisternum
- At 32nd week : At two-thirds of the distance between the umbilicus and xiphisternum
- At 36th week : At the level of xiphisternum
- At 40th week : Sinks back to the level of the 32 nd week, but the flanks are full, unlike that in the 32 nd week



COUNSELLING IS AN IMPORTANT ADJUNCT TO MANAGEMENT

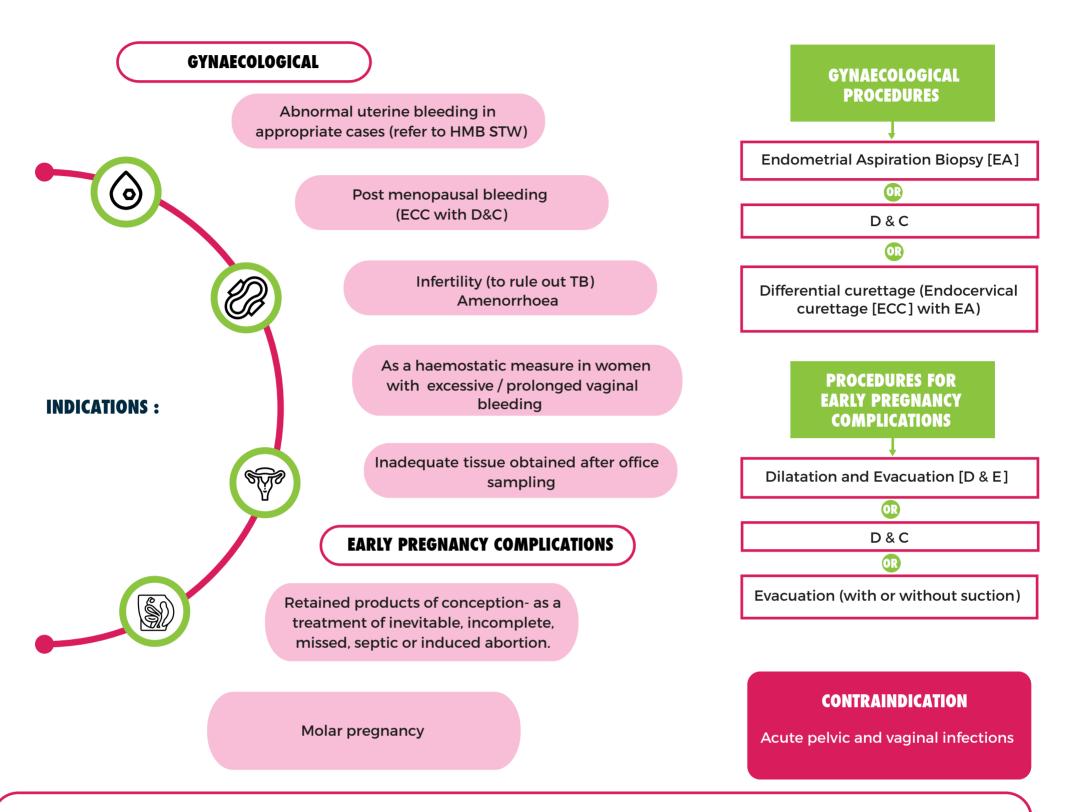
🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES





Standard Treatment Workflow (STW) for DILATATION AND CURETTAGE (D&C)

- Mostly done for gynaecological indications, but may also be considered in early pregnancy complications
- Though office endometrial biopsy using either thin flexible or Karman cannula or office hysteroscopy has obviated the need for traditional D&C in gynaecological cases, it still has a place when other modalities are not available or do not yield adequate tissue



WHERE CAN IT BE PERFORMED?

- In secondary or tertiary healthcare centres preferably where facilities for anaesthesia and operation theatre are available to deal with procedure related complications, if any.
- Endometrial aspiration biopsy is usually done as an outpatient procedure in non pregnant cases.

ALL TISSUE REMOVED MUST BE SENT FOR HISTOPATHOLOGICAL EXAMINATION

PRE- OPERATIVE REQUISITES							
Presence of a valid indication General medical fitness & no contraindication A written informed conservation							
ANESTHESIA (ANY OF THE FOLLOWING)							
 General anesthesia Regional a 	\cdot General anesthesia \cdot Regional anesthesia \cdot Paracervical block with 1% xylocaine \cdot IV sedation \cdot IM/ oral analgesia						
Strict asepsis to be maintained. Antibiotics to be used judiciously and decided as per need of individual case.							
 POST PROCEDURE CARE & FOLLOW UP Observe the patient for minimum two hours after the procedure for haemorrhage or any other symptoms or signs of complications prior to discharge Patient can be discharged as soon as she is 	COMPLICATIONS • Excessive bleeding • Cervical laceration • Perforation of the uterus • Injury to bowel and bladder	DO • Evacuation of uri before procedure • Safety checklist • Dorsal/lithotomy • Bimanual pelvic	inary bladder e. / position	DONT'S Over abduction of legs No sounding in cases of pregnant uterus. No forceful insertion of any instrument 			

- · Patient can be discharged as soon as she is comfortable and alert.
- Most common side effect is abdominal cramps which can be managed by oral analgesics.
- Warning signals to report backere to be explained at the time of discharge - severe pain, bleeding, foul smelling discharge or fever.
- Follow up is done after a week with histopathology report for further advice.
- Pelvic infection
- Post-operative intra uterine adhesions
- Bimanual pelvic examination prior to the procedure
- Sounding to measure
- uterocervical length ONLY in non pregnant women.
- Sample to be sent for histopathology and microbiology (where indicated)
- REFER in case of a complication
- any instrument
- Abandon the
- procedure in case of suspected perforation and refer to higher centre.
- Insertion of the dilator should be just beyond the internal os and NOT till the fundus

D&C is a blind procedure and may miss the pathology in some cases. In cases where focal pathology is suspected, tissue should be obtained under hysteroscopic visualization.

COUNSELLING IS AN IMPORTANT ADJUNCT TO MANAGEMENT

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



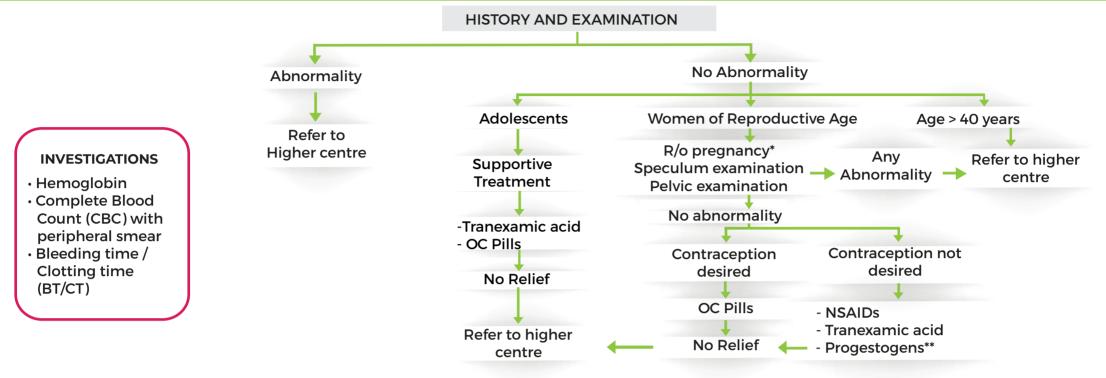


NDIAN COUNCIL MEDICAL RESEAR Serving the nation since

Standard Treatment Workflow (STW) for the Management of HEAVY MENSTRUAL BLEEDING (HMB)

ICD-10-H90.5

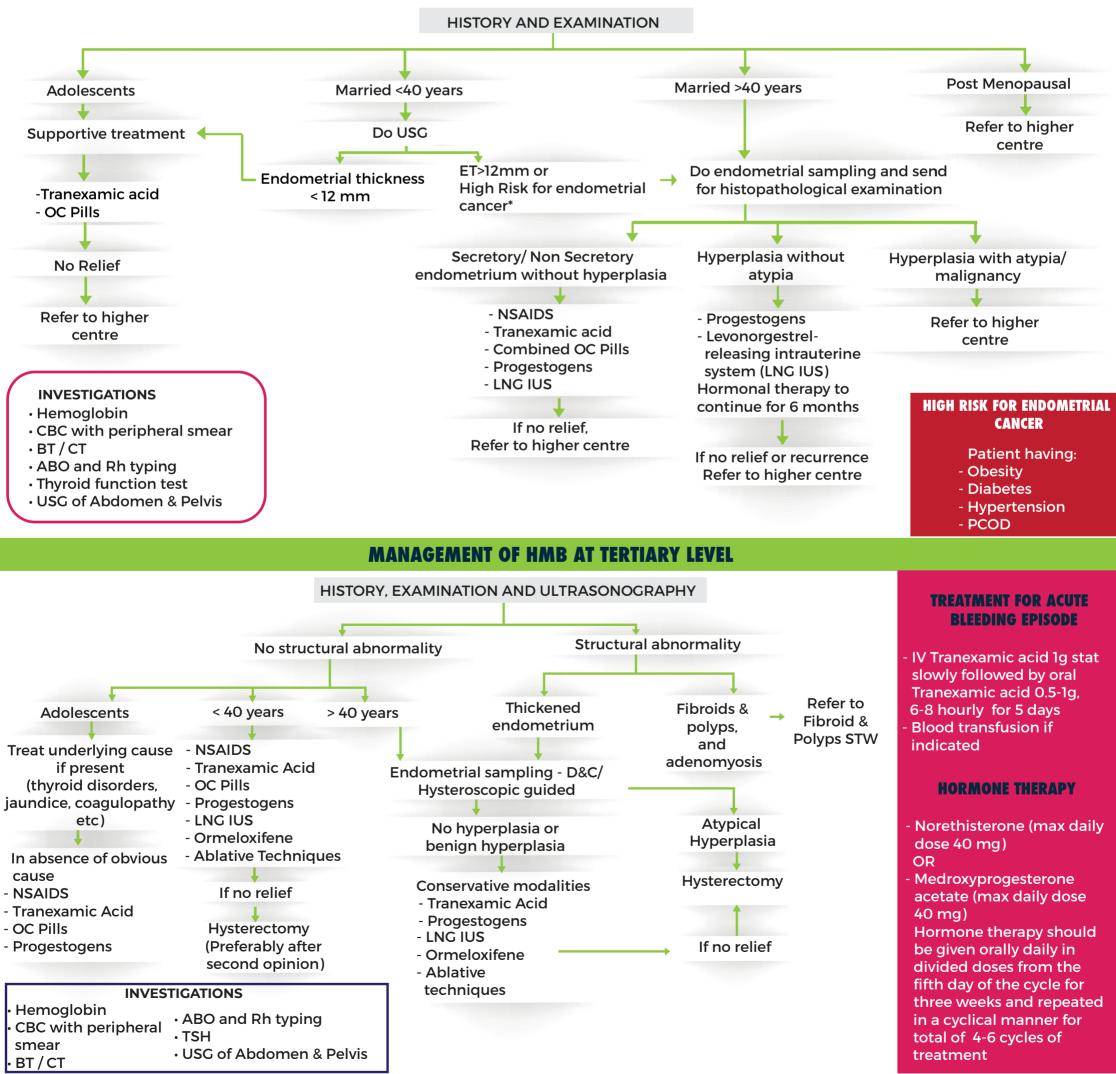
TO DO AT ALL LEVELS HISTORY **EXAMINATION** SUPPORTIVE TREATMENT • General • Age **Evaluate pallor** Parity **Calculate BMI** Reassurance · Detailed menstrual history including irregularities Hematinics · Other medical illness: thyroid disorder, Systemic Tranexamic acid during episode of coagulopathy, jaundice etc CVS, RS and hepatosplenomegaly heavy bleeding IUCD use Lactation Local examination (where indicated and Drug intake feasible) P/S and P/V MANAGEMENT OF HMB AT PRIMARY LEVEL



* R/o Pregnancy in doubt especially in all women of reproductive age group after appropriate consent

** Amongst progestogens Norethisterone provides the best hemostasis

MANAGEMENT OF HMB AT SECONDARY LEVEL (CHC)



COUNSELLING IS AN IMPORTANT ADJUNCT TO MANAGEMENT

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



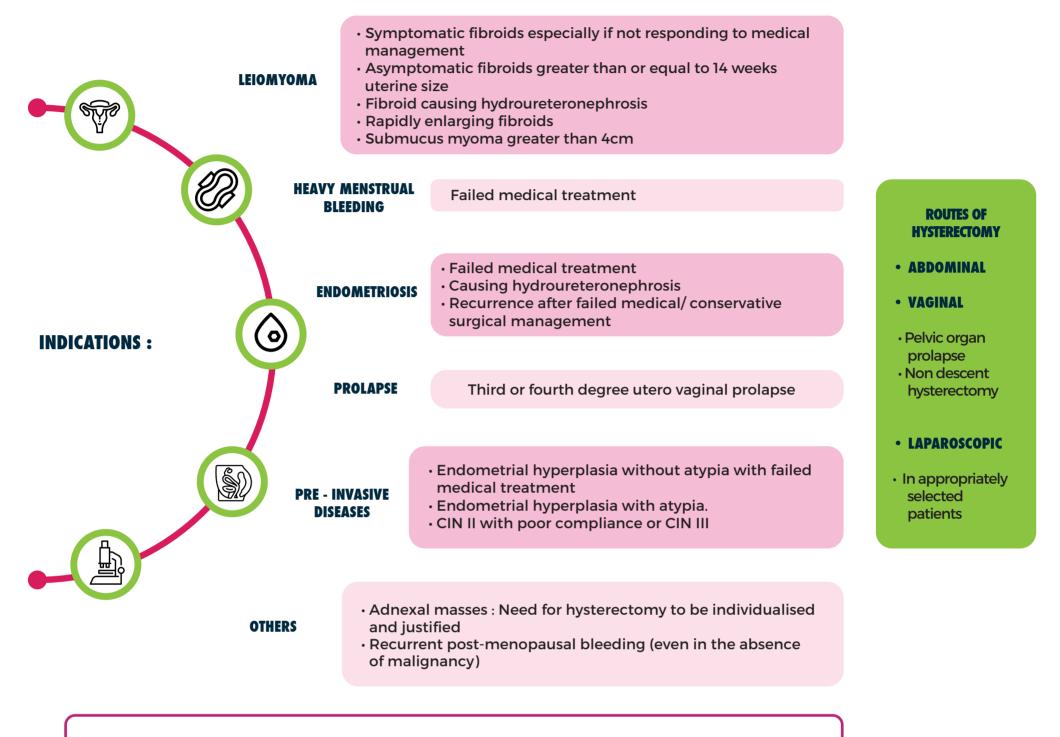




Standard Treatment Workflow (STW) for HYSTERECTOMY FOR BENIGN GYNAECOLOGICAL CONDITIONS

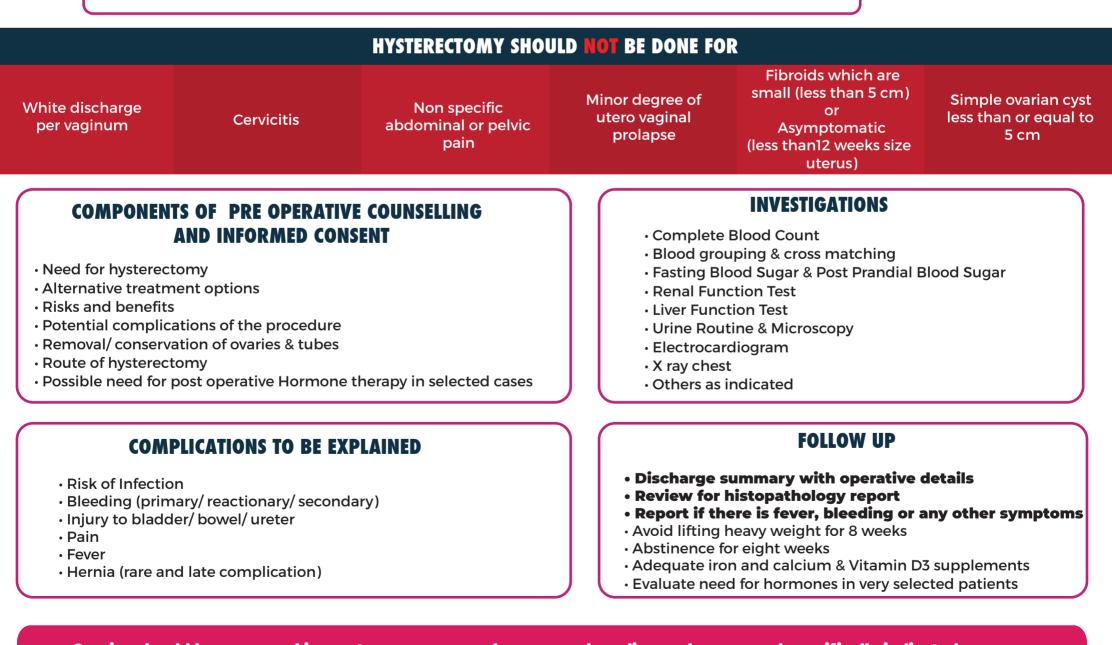
IN WOMEN AGED LESS THAN 40 AND/OR LOW PARITY IT IS MANDATORY TO HAVE A SECOND OPINION FROM A QUALIFIED GYNAECOLOGIST

HYSTERECTOMY TO BE CONSIDERED ONLY WHEN CHILD BEARING IS COMPLETED & RARELY IN YOUNGER PATIENTS



Simple ovarian cysts less than 5 cm in size and without other significant/ suspicious features





Ovaries should be preserved in most pre-menopausal women unless diseased or removal specifically indicated
 While doing hysterectomy for benign gynaecological conditions in pre-menopausal women, it is recommended to combine it with bilateral salpingectomy with a view to minimise the risk of subsequent development of ovarian malignancy ¹²

1. Pérez-López FR et al, Interventions to reduce the risk of ovarian and fallopian tube cancer: A European Menopause and Andropause Society Postition Statement. Maturitas. 2017

2. Darelius A et al, Efficacy of salpingectomy at hysterectomy to reduce the risk of epithelial ovarian cancer: a systematic review. BJOG. 2017.

COUNSELLING IS AN IMPORTANT ADJUNCT TO MANAGEMENT

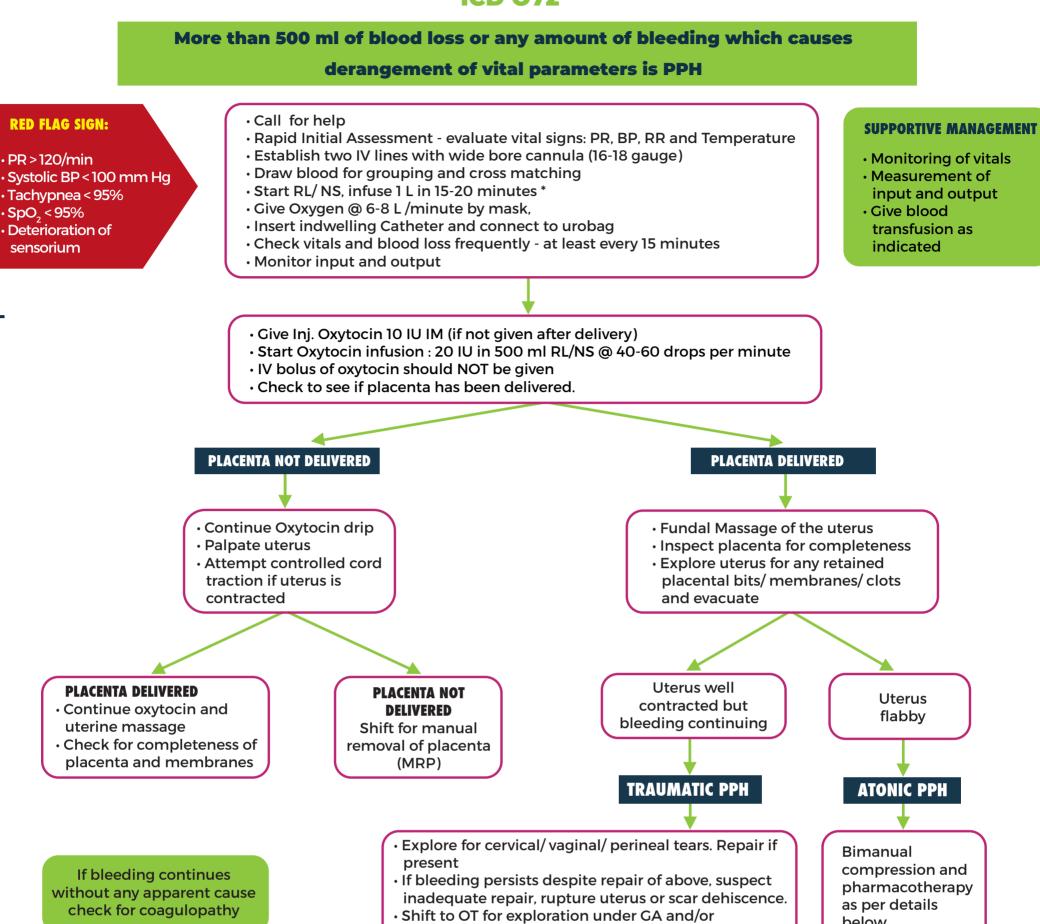
KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES





Standard Treatment Workflow (STW) for the Management of **POSTPARTUM HAEMORRHAGE (PPH)**

ICD 072



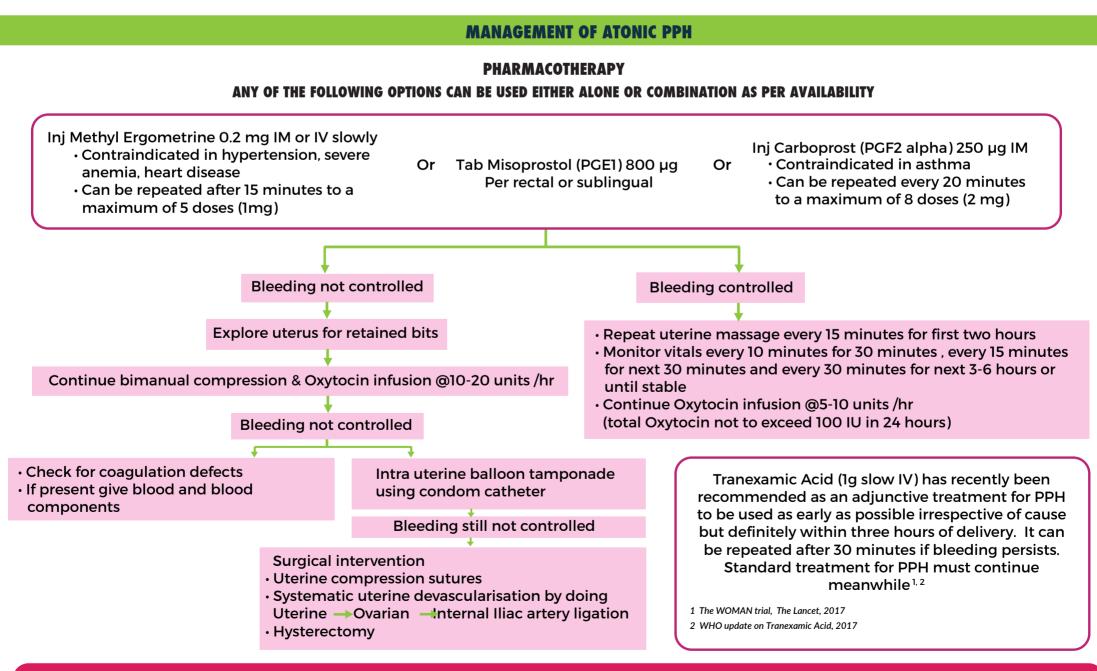


laparotomy

below

* Arrange for blood / blood product at the earliest

3 ml of crystalloid solution should be used to replace every ml of blood lost during the initial part of the acute bleeding phase



Timely Referral to a higher centre must be considered if facilities for blood transfusion or exploration and surgical intervention are not available. Patient must be transported with I/V fluids containing oxytocin on flow and preferably with uterine/vaginal tamponade in situ.

- Aortic compression may be used as a short time measure to reduce blood loss while awaiting definitive steps.
- Non- pneumatic anti-shock garment (NASG) should be used during transport if available
- Uterine artery embolization may be offered in selected patients if facilities are available

COUNSELLING IS AN IMPORTANT ADJUNCT TO MANAGEMENT

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

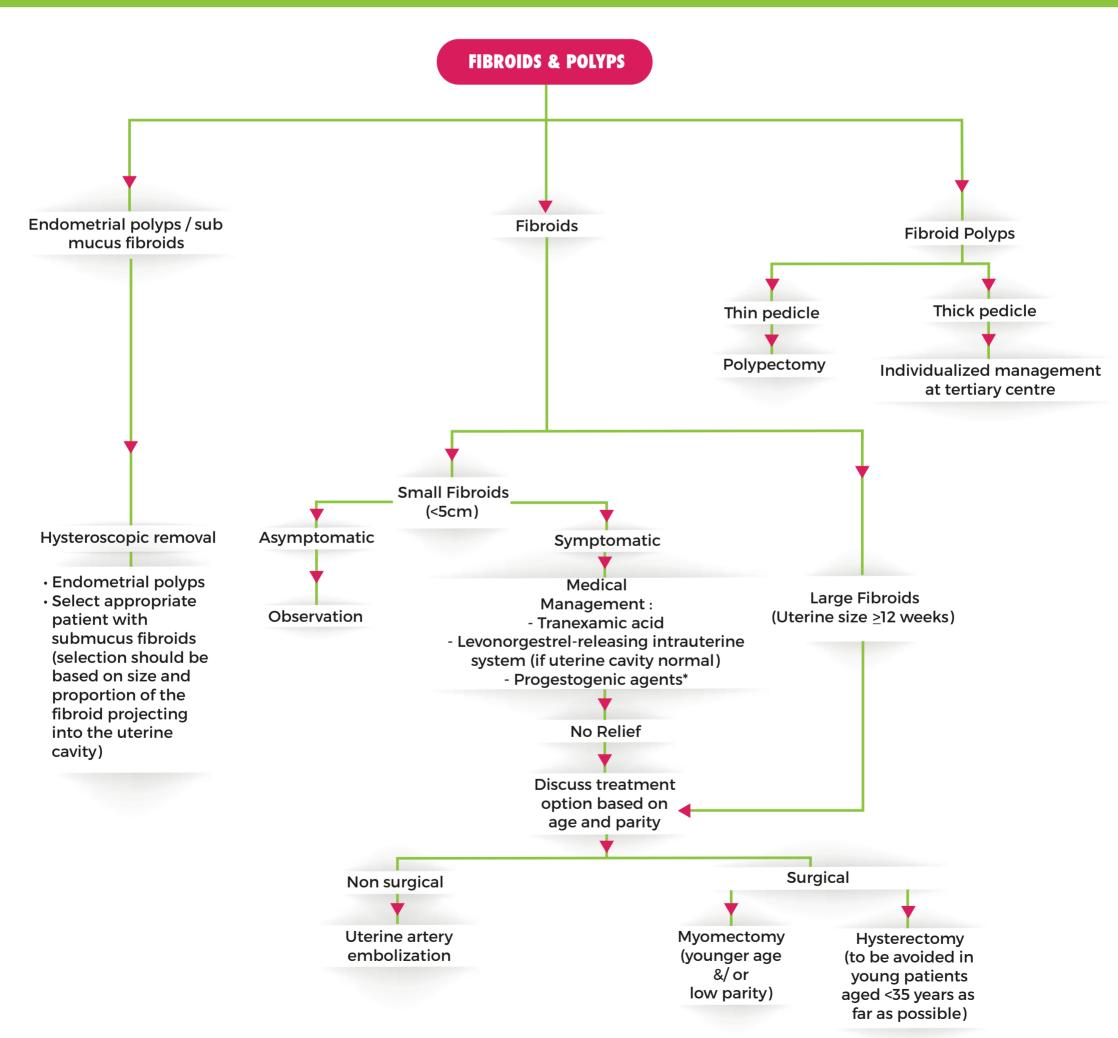




Standard Treatment Workflow (STW) for the Management of UTERINE FIBROIDS AND POLYPS ICD-10-D25 & N84

Heavy menstrual/irregular **EXAMINATION** bleeding **GPE**: Specially check for pallor Abdominal examination: Check for any Urinary pressure symptoms suprapubic mass or lump **P/S**: Inspect cervix for local abnormalities Heaviness and pain in P/V: Assess for uterine size (enlarged abdomen, dysmenorrhoea and/or irregular uterus) S) **SYMPTOMS:** Mass in lower abdomen INVESTIGATIONS Hemoglobin Infertility **Complete blood count** Thyroid function test Asymptomatic 2141a USG

ASYMPTOMATIC FIBROIDS <5CM DO NOT NEED TO BE TREATED



*Norethisterone (max daily dose 40 mg) OR Medroxyprogesterone acetate (max daily dose 40 mg). Any hormone should be given orally daily in divided doses for a duration of three weeks and repeated in a cyclical manner for total of 4-6 cycles of treatment

ALL THERAPUTIC OPTIONS NEED TO BE EXPLAINED TO THE PATIENT INCLUDING JUST KEEPING THE PATIENT ON OBSERVATION. ALL PATIENTS OF FIBROID UTERUS DO NOT NECESSARILY NEED HYSTERECTOMY.

COUNSELLING IS AN IMPORTANT ADJUNCT TO MANAGEMENT

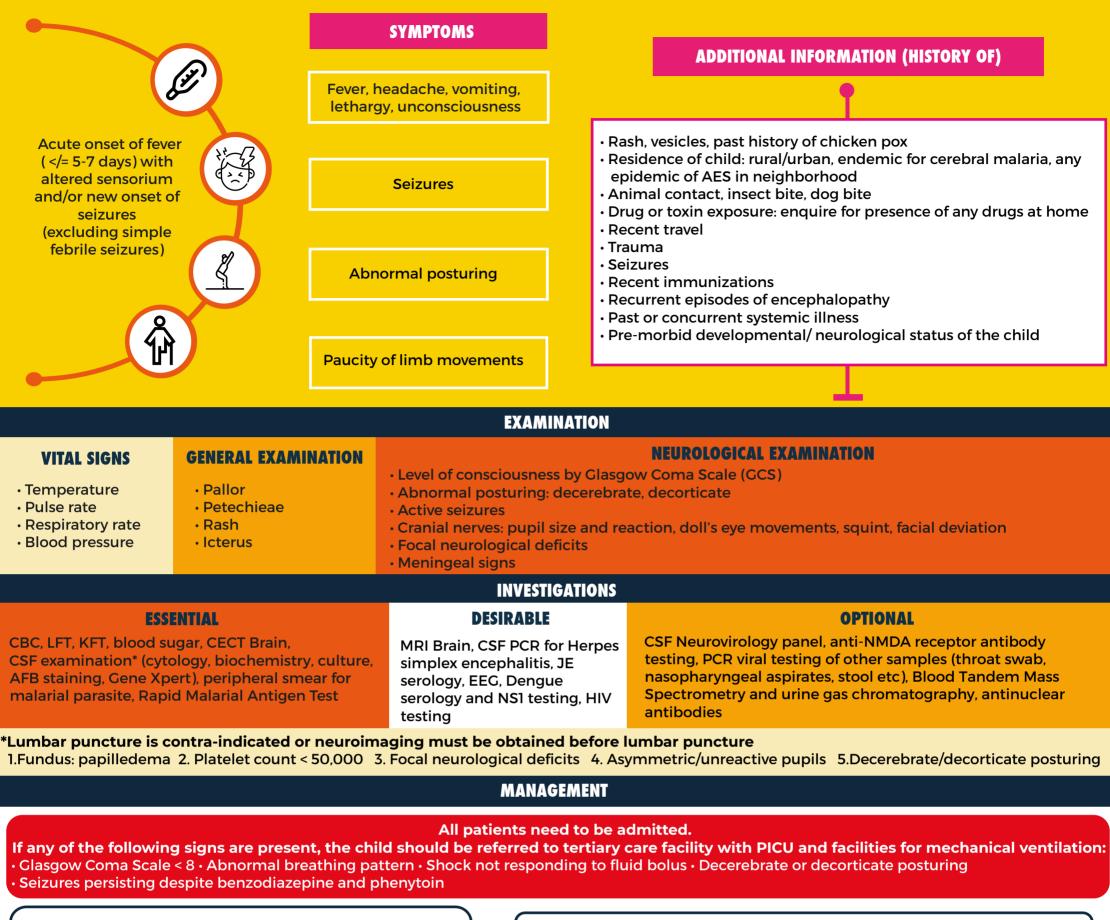
KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



PAED ATRICS



Standard Treatment Workflow (STW) for the Management of **ACUTE ENCEPHALITIS SYNDROME (AES) IN CHILDREN ICD-10-G04**



Step I: Rapid assessment and stabilization Establish and maintain airway: Intubate if GCS<8.

Step II: History, Examination and Investigations as given above

- impaired airway reflexes, abnormal respiratory pattern, signs of raised intracranial pressure, SpO2 <92% despite high flow oxygen and fluid refractory shock
- Ventilation, oxygenation
- Circulation: Establish IV access, take samples for relevant investigations, fluid bolus if in circulatory failure (20 mL/kg NS), inotropes if required
- · Identify signs of cerebral herniation or raised ICP
- Temperature: treat fever and hypothermia
- Treat ongoing seizures- Benzodiazepine, followed by phenytoin loading

Step V: Prevention/treatment of complications and rehabilitation

- Physiotherapy, posture change, prevent bed sores and exposure keratitis
- Complications: aspiration pneumonia, nosocomial infections, coagulation disturbances
- Nutrition: early feeding
- Psychological support to patient and family

#Management of raised intracranial pressure

Step III: Empirical Treatment (must be started if CSF cannot be done/ report will take time and patient sick)

- Ceftriaxone: 100 mg/kg/day in 2 divided doses X 10-14 days
- Acyclovir (use in all suspected sporadic viral encephalitis):
- 3 mo to 12 y: 500mg/m2 8 hourly (min 21 days)
- >12 y: 10mg/Kg 8 hourly (14-21 days in confirmed cases)**
- Artesunate combination therapy (stop if peripheral smear and RDT are negative): 3mg/kg in child <20 kg, and 2.4mg/kg in child > 20kg IV/IM at 0.12 and 24 hours, followed by once daily parental/oral X 3-7 days

**If therapy was started empirically stop acyclovir, in case an alternative diagnosis is confirmed, or HSV PCR of CSF is negative on two occasions (24-48 h apart) and MRI imaging not suggestive of Herpes Simplex Encephalitis

Step IV: Supportive care and treatment

- Maintain euglycemia, hydration and control fever
- Treat raised intracranial pressure#, mild head-end elevation-15-30°
- Treat seizures##: Give anticonvulsant if: history of seizures / GCS <8 / child has features of raised ICP
- Steroids: Pulse steroids (methylprednisolone) to be given in children with suspected acute disseminated encephalomyelitis or autoimmune encephalitis

##Treatment of seizures

- Intubate if: GCS <8 / evidence of herniation / irregular respirations and inability to maintain airway
- Signs of impending herniation: patient to be hyperventilated to a target PaCO2 of 30-35 mmHg
- Initial bolus of Mannitol(0.25 g/kg), then 0.25 g/kg q 6 h as per requirement, up to 48 hours.
- In the presence of hypotension, hypovolemia, and renal failure: hypertonic (3%) saline (preferable to mannitol) 0.1-1 mL/kg/hr by infusion; serum sodium to be targeted to 145-155 meg/L
- Adequate sedation and analgesia
- Avoid noxious stimuli
- Administer pebulized lignocaine prior to endotracheal tube suction

1st Line: IV Lorazepam 0.1mg/kg or Midazolam 0.2 mg/kg
orDiazepam 0.3 mg/kg).
If no IV access: IM Midazolam 0.2 mg/kg
2nd Line: Inj. Phenytoin 20 mg/kg (in Normal saline
Img/kg/min)
If seizures still persist:
Refractory status: Transfer to PICU -> midazolam infusion
(1-18 microgram/kg/min)
If ICU facilities not available: sodium valproate (20 mg/kg) or
levetiracetam (20-40 mg/kg) or phenobarbitone (20mg/kg)

Administer hebdlized i	ignocame phor to endo	liacheal tube such	lorning		
			DISCHARGE CRITERIA		
Hemodynamically stable	Improvement in consciousness	Afebrile	Has started eating and drinking orally	Seizures have subsided	Parents have been explained the supportive care and physiotherapy to be continued at home
		• • • • • • • • • • • • • • • • • • •			

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

REFERENCES

- 1. World Health Oraganisation. Acute Encephalitis Syndrome. Japanese encephalitis surveillance standards. January 2006. From WHO-recommended standards for surveillance of selected vaccine-preventable diseases. WHO/V&B/03.01. Available from: http://www.who.int/vaccines-documents/ DocsPDF06/843.pdf
- 2. National Program for Prevention and Control of Japanese Encephalitis/Acute Encephalitis Syndrome 2014. Government of India Ministry of Health & Family Welfare Directorate General of Health Services National Vector Borne Disease Control Programme.
- 3. Sharma S, Mishra D, Aneja S, Kumar R, Jain A, Vashishtha VM. Consensus guidelines on evaluation and management of suspected acute viral encephalitis in children in India. Indian Pediatr. Nov 2012;49(11):897-910. 4. Sankhyan N, Vykunta Raju KN, Sharma S, Gulati S. Management of raised intracranial pressure. Indian J Pediatr. 2010 Dec;77(12):1409-16.





Standard Treatment Workflow (STW) for the Management of **ACUTE DIARRHEA** ICD-10-R19.7

 DIARRHEA IS >3 loose or watery stools/ day Acute Diarrhea <14 days Persistent diarrhea >14 days Dysentery - blood in stools Que to to to to to to to to to to to to to	 REFER TO HOSPITAL Severe malnutrition/HIV Severe dehydration Hypernatremic (Na >145mmol/L)/ hyponatremic dehydration (Na <135 mmol/L) Dysentery with age <1 yr/ measles in past 6 weeks/ dehydration/ sick Dysentery with no improvement on antibiotics Persistent diarrhea with dehydration Persistent diarrhea with serious systemic infection such as pneumonia, sepsis, infants <4 months of age, or when there is no improvement with treatment over 5 days 		
		MANAGEMENT SSIFY DEHYDRATION	
Not enough signs to classify some or severe dehydration	2 of following a) Restless , in b) Sunken eye c) Drinks eage d) Skin pinch	ritable es	2 of following: a) Lethargy / unconscious b) Sunken eyes c) Not able to drink/ drinking poorly d) Skin pinch - goes back slowly
 NO DEHYDRATION: PLAN A Fluids Give extra fluids (as much as a will take) until diarrhea stops. Use WHO ORS after each loos stool (in addition to usual fluid intake) Upto 2 yrs → 50 -100 ml 2 yrs or more → 100 -200m On ORS packet check wheth 200ml or 1 litre of clean water i needed Frequent small sips with spoor cup. If child vomits, wait 10 minute then continue slowly. Homemade fluids- salted rice water, salted yogurt drink, vegetable or chicken soup with and clean water unsweeteneous 	 Manage in clinic /day amount of ORS (75m) If weight is not known If weight is not known If weight is not known Mage AGE <	11 ths12 - 23 months $2 - 4$ years $5 - 14$ years15 years or older $(.9)$ $8 - 10.9$ kg $11 -$ 15.9 kg $16 -$ 29.9 kg 30 kg or more $(.0)$ $600 -$ 800 $800 -$ 1200 $1200 -$ 2200 $2200 -$ 4000 $(.10)$ $600 -$ 800 $1200 -$ $1200 2200 -$ 4000 $(.10)$ $600 -$ 800 $1200 -$ $2200 2200 -$ 4000 $(.10)$	SEVERE DEHYDRATION: PLAN C 9. Or the second seco
and clean water, unsweetenec fruit juice and coconut water	• Danger signs*		 If child can drink, give ORS by mouth while

- fruit juice and coconut water Unsuitable fluids - carbonated
- · Hand washing , proper disposal of excreta
- beverages, commercial fruit juice, sweetened tea & coffee, other medicinal teas / infusions.
- Zinc supplement (Zinc sulphate/ carbonate / acetate)
 - \cdot 2-6 months \rightarrow 10mg/day x 2 weeks
 - \cdot >6 months \rightarrow 20mg/day x 2 weeks

Counsel Mother/ Attender

Feeding advise

- Infants on breast feed, to continue more frequent breast feeding than usual.
- Those not on breast feed to continue their usual milk feed/ formula at least once in 3 hours.
- Give age appropriate foods to >6 months old based on their pre illness feeding pattern

Danger signs (return immediately)

- Passing many watery stools
- Repeated vomitings / very thirsty
- Eating / drinking poorly
- Develops fever / blood in stools

Follow up in 5 days if no improvement

Safe drinking water

Hygiene practices

- Appropriate feeding practices
- Vaccination as per IAP guidelines

INVESTIGATIONS

Some dehydration:

- **Preferable Tests-** electrolytes
- Severe dehydration:
- **Essential tests-** CBC, electrolytes Preferable Tests- Renal Function Tests, VBG
- In suspected cholera cases:
- Preferable tests- stool for hanging drop and stool culture
- Dysentery: (no response to antibiotic in 2 days) Preferable test- stool culture & stool routine for trophozoites of Ameoba Persistent diarrhea:
- Preferable test- stool routine microscopy,
- urine routine microscopy, urine culture, sepsis screen

WHEN CONSIDERING ALTERNATIVE DIAGNOSIS OF **PERSISTENT DIARRHEA AND DYSENTRY**

- **PERSISTENT DIARRHEA**
- Appropriate fluids to prevent or treat dehydration Nutrition:
 - · If breastfeeding, give more frequent, longer breastfeeds, day and night. Other milk: replace with increased breastfeeding, or with fermented milk products, such as yogurt, or half the milk with nutrient-rich semi-solid food.
 - For other foods, follow feeding recommendations for the child's age: give small, frequent meals (at least 6 times a day), and avoid very sweet foods or drinks.
- Zinc for 14 days
- Supplement vitamins / minerals
- Antimicrobial to treat diagnosed infection
 - A) Intestinal infection:
 - If blood in stool: Treat like dysentery
 - If stool routine suggestive of Amoebiasis: Treat for it
 - If stool suggestive of cyst/Trophozoite of Giardia: Give Metronidazole
 - 5mg/kg/dose x 8hrly x 5 -7 days B) Treat Non intestinal such as UTI / Otitis Media
- Follow up in 5 days
- Refer to hospital (See box)

REFERENCES

- 1. IMCI (WHO) module on Diarrhea 2014.
- 2. WHO Treatment for Diarrhea A manual for physicians and other senior health workers 2005.
- 3. WHO GLOBAL TASK FORCE ON CHOLERA CONTROL 2010.

🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

This STW has been prepared by national experts of India with feasibility considerations for various levels of healthcare system in the country. These broad guidelines are advisory, and are based on expert opinions and available scientific evidence. There may be variations in the management of an individual patient based on his/her specific condition, as decided by the treating physician. There will be no indemnity for direct or indirect consequences. Kindly visit our web portal (stw.icmr.org.in) for more information. © Indian Council of Medical Research and Department of Health Research, Ministry of Health & Family Welfare, Government of India.

- Assess heart rate/respiratory rate/BP/CFT/ consciousness and recognize early shock
- Refer for hospitalization

the drip is set up

 If prevalance of cholera – Doxycycline single dose 300mg or Tetracycline 12.5mg/kg 4 times a day x 3 days. For young children Erythromycin 12.5mg/kg 4 times a day x 3 days

- Associated vomitings Ondanstetron 0.15 mg/kg/dose IV/oral in addition to rehydration therapy
- Reassess every 15-30 minutes till a strong radial pulse is present and then every hour If hydration status is not improving, give IV drip more rapidly
- After 6 hours (infants) and 3 hours (older patients) - evaluate for dehydration and choose the appropriate plan (A, B, or C) to continue treatment
- Give ORS (about 5 ml/kg/hour) as soon as the child can drink: usually after 3-4 hours (infants) or 1-2 hours (children)
- Observe for 6 hours after the child has been fully rehydrated.
- In hypernatremic and hyponatremic dehydration child appears relatively less ill / more ill respectively and needs to be referred for hospitalization

DISCHARGE CRITERIA

- Suffcient rehydration (indicated by wt gain &/or clinical status)
- IV fluids no longer needed
- Oral intake = /> losses
- Medical f/u available

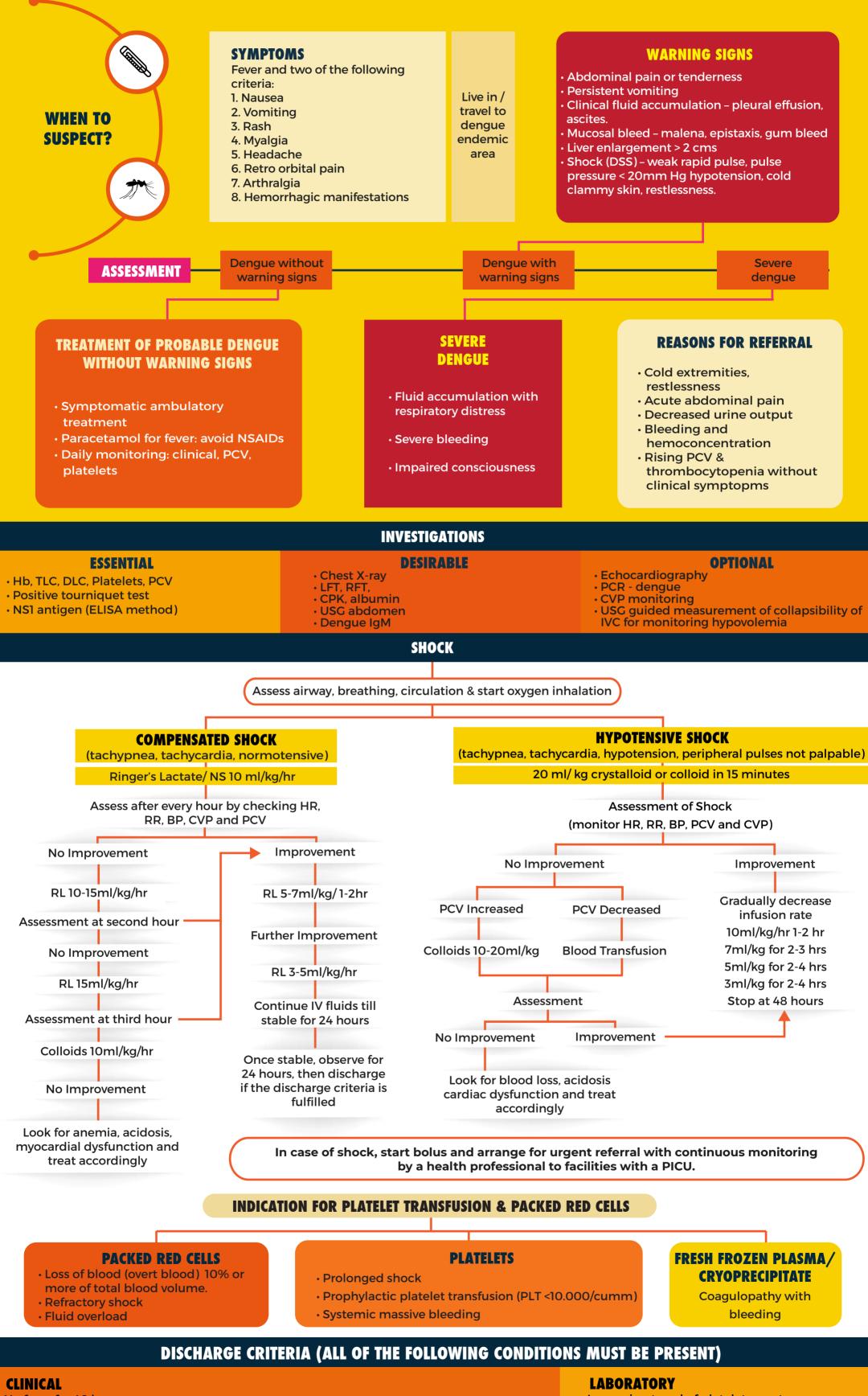
DYSENTERY

- Treat dehydration according to assessment.
- Ciprofloxacin 15mg/kg twice a day and reassess after 2 days.
 - Improvement: 3 days of treatment
- No improvement \rightarrow Cefixime 10 mg//kg/d, 2 div doses. Reassess after 2 days. If better complete 3 -5 days of treatment.
 - If stool routine positive for Ameobiasis : Metronidazole 10mg/kg/dose 8 hourly x 7 days (10 days in severe cases)
- Refer to hospital (See box)





Standard Treatment Workflow (STW) for the Management of DENGUE FEVER ICD-10-A90



No fever for 48 hours

• Improvement in clinical status (check for general well-being, appetite, haemodynamic status, urine output, respiratory distress)

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

- Increasing trend of platelet count
- Stable haematocrit without intravenous fluids





Standard Treatment Workflow (STW) for the Management of **FEVER IN CHILDREN ICD-10-R50**

FEVER IS Core (rectal) temperature $\ge 38.0^{\circ}$ C (100.4°F) or axillary temperature $> 37.5^{\circ}$ C (100.4°F).

		EXAM		CLUES TO A SPECIFIC DIAGNOSIS		
	Fever duration	Vital signs: Temp, H		Fever + respiratory symptoms: Cough, runny nose: URTI Membrane over tonsils/pharynx:		
	Localizing symptoms of: RTI, UTI, GI tract infection, CNS infection	Appearance: Sick, to inconsolable, dehydra	oxic, lethargic, irritable,	Diphtheria Paroxysmal cough: Pertussis like illness Barking cough: Laryngotracheobronchitis/ croup		
WHAT TO ASK?	Rash, joint symptoms, skin/soft tissue swelling or redness Vaccination within 24 hours,	General Examination: • Ear, nose, throat • Rash (petechieae, macules, papules, vesicles, nodules, polymorphic) • Lymphadenopathy • Skin (pustules, pyoderma, impetigo, cellulitis) • Joints		 Fever + rash Red maculopapular rash: Measles, Rubella, Dengue. Fine generalized maculopapular rash with systemic dysfunction/shock: Meningococcemia. Itchy erythematous macules evolving to clear vesicles: Varicella 		
Family/ neighbourhood history of similar illness		 Genitalia (for erythema, tenderness, edema) Bones Systemic Examination Chest auscultation, abdominal palpation, CNS, CVS 		 Fever + other symptoms: Parotid gland swelling: Mumps Arthritis: Consider Chikungunya, acute rheumatic fever, JIA Strawberry red tongue, skin peeling, lymphadenopathy, conjunctival injection: Kawasaki disease 		
INVESTIGATION OF THE FEBRILE CHILD (Consider if one or more of the following are warranted. Perform investigations only where result impacts management)						
<7 DAYS FEVER ALON	<pre><7 DAYS FEVER ALONE </pre> <7 DAYS AND LOCALIZING SYMPTOMS PRESENT <7 DAYS AND NON SPECIFIC SYMPTOMS >7 DAYS AND FEVER ALONE SYMPTOMS >7 DAYS AND LOCALIZING SYMPTOMS >7 DAYS AND LOCALIZING SYMPTOMS					

ESSENTIAL:

If fever <72 hours and child not looking sick: No investigations If fever >72 hours, consider: TLC, DLC, P.S for leukocyte morphology, malarial parasite & platelet count

DESIRABLE: Rapid antigen test for malaria, NS1 antigen and dengue IgM antibody, blood culture

ESSENTIAL: As given in the

DESIRABLE: As given in

the first box + consider:

microscopy & culture,

first box + consider:

aspiration.

chest Xray, CSF analysis

OPTIONAL: As given in the

ultrasonography, throat/

pharyngeal swab, pus

(Clean-catch) urine

first box

ESSENTIAL: As given before

DESIRABLE: As given before. Additionally consider: serology for specific viral infection, rapid antigen test for malaria, NS1 antigen and dengue IgM antibody, blood culture, serology for scrub typhus

OPTIONAL: As given before

SYMPTOMS

ESSENTIAL: All mentioned in Essential & Desirable list in the prior boxes. Additionally consider Widal test.

DESIRABLE: Consider Mantoux test, ultrasonography

OPTIONAL: As given before. Additionally consider: Ultrasonography of abdomen, chest, pericardium, joint(s), abscess, lymph node clusters, parotid gland etc, for microscopy, Xpert MTB RIF assay, Mycobacterial culture. Consider:

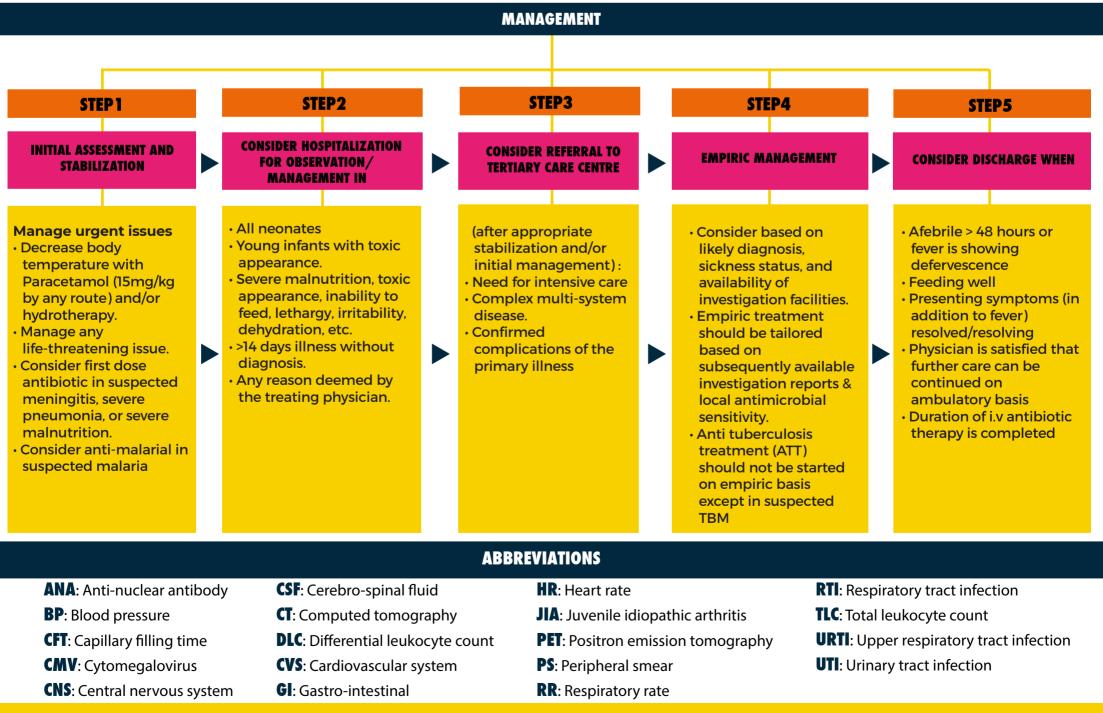
ESSENTIAL: All investigations mentioned in the prior boxes

DESIRABLE: All

investigations mentioned in the prior boxes. Additionally consider: serology for Brucella, CMV, Herpes, Japanese encephalitis. CT scan in deep seated abscess or lung abscess, Bone marrow examination, ANA profile, HIV serology, PET scan.

OPTIONAL: All





KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

REFERENCES

- 1. World Health Organization. Integrated Management of Childhood Illness: distance learning course.
- http://apps.who.int/iris/bitstream/handle/10665/104772/9789241506823_Module-5_eng.pdf;jsessionid=942F89F89671BA396EC7F46C9B5C1158?sequence=7
- 2. Mahajan P, et al. Consensus Guidelines on Evaluation and Management of the Febrile Child Presenting to the Emergency Department in India. Indian Pediatr 2017; 54: 652-60.
- 3. World Health Organization 2015. Government of India National Guidelines for Clinical Management of Dengue Fever.
- 4. Kliegman RM (ed). Nelson Textbook of Pediatrics 20th edition, 2016.







Standard Treatment Workflow (STW) for the Management of SEPSIS AND SEPTIC SHOCK IN CHILDREN

ICD-A41.9, R65.21

		WHE	IN TO SUSPECT	r (2-59 MONT	HS)?	CHECK FOR HISTORY OF
Sepsis to be suspected:		Poor Feeding	Lethargy	Decreased	Unconsciousness	Prior treatment
in children with any infections (fever with		recarry		responsiveness		Previous recurrent infections
or without rashes/ pneumonia/	ANN ANN	Cold/ bluish peripheries	Rapid or shallow breathing	Chest in drawing	Stridor	Prior hospitalisation
diarrhoea) and they are at risk of life			Convulsions	Stiff neck	Chronic systemic illness (congenital or acquired)	
threatening organ dysfunction		vomiting	urine output	Convuisions	Still Heck	Immunization (age appropriate)

EXAMINATION

GENERAL PHYSIC	L EXAMINATION	VITAL SIG	SYSTEMIC EXAMINATION		
Lethargy Decreased alertness	Petechial rash Mucosal bleeding	Pulse volume (High volume as well as low volume/feeble pulse)	Heart rate and respiratory rate (outside the age range)	Respiratory: Signs of respiratory distress - retraction, nasal flaring, grunting ,crepitation on auscultation	
Activity	Rapid breathing	Capillary refilling time > 3	Pulse oximetry	CVS: Murmur, gallop rhythm Per abdomen: Abdominal distension	
Pallor	Chest in drawing	seconds	(saturation <95%)	CNS : *AVPU scale, signs of meningitis,	
Cyanosis	Cold peripheries	Blood pressure* (Systolic blood Pressure	>1 year child if systolic BP < 70+ Age	seizures Skin : Rashes	
Skin mottling	Assess nutritional status	< 70 in <1 year)	(yrs) x2) or (lower than age range)	Bone & joints: Swelling, redness, tenderness	

SIGNS OF SEVERE DEHYDRATION

Diarrhoea plus any two of these: Lethargy or unconscious, not able to drink or drinks poorly, Sunken eyes, skin pinch goes back very slowly

INVESTIGATIONS- (Based on symptoms and available facility)

Essential – Complete blood counts, peripheral blood film, urine routine, blood sugar, CRP, serum electrolytes, renal function test, liver function test **Desirable** - Blood culture, blood gas, relevant cultures (based on symptoms), chest X-ray, specific illness- Malaria – rapid malarial antigen test, Dengue- dengue NS1, IgM, CSF study

Optional- PCT , USG to guide the fluids

MANAGEMENT

DIAGNOSTIC ALGORITHM

CHILD (2-59 MONTHS OF AGE WITH FEBRILE ILLNESS (WITH WARNING SIGNS)

GOOD PERIPHERAL PERFUSION

Admit or initiate treatment as per IMNCI guidelines²

With fast pulse, cold peripheries, poor pulse volume, CRT >3 seconds (Fast pulse: HR> 180 in < 12 month old child, HR >120 in >12 month old child)

POOR PERIPHERAL PERFUSION**

**If there is improvement after 1st bolus and history of diarrhea present then:

Give 70 ml/kg over 5 hours in infants and over 2 ½ hours in a child with hypovolemic shock. Give additional fluids if losses continue.

Start maintenance fluid in case of other illness

Antibiotics

- 1. >3 months Inj Ceftriaxone 100mg/kg/day (2 divided doses)
- 2. <3 month Inj Cefotaxime 200mg/kg (divided 6-8hrly),
 - Inj Gentamicin 5-7.5 mg/kg single dose /day
- 3. If soft tissue infection: consider Inj Cloxacillin 200mg/kg divided
 6 hourly or Inj Amoxicillin- Clavulanic acid 30 mg/kg/dose 8hrly)

Inj Adrenaline- 0.3x body weight in mg in 50 ml NS or 5% dextrose at 1 ml/hr will give 0.1 microgram/kg/min

Admit, initiate treatment, refer to centre with facility of ICU, ventilation, 24 hour monitoring (if required)

Start O₂ with face mask @ 4-6 lit/min, or hood @8-10 lit if not available nasal prongs 1-2 lit/min to maintain SpO₂ >95%, Insert two IV cannulas, give first dose of antibiotics within first one hour

Give 20 ml/kg of normal saline fluid bolus over 20- 30 minutes.

Reassess for decreases in heart rate, improvement in pulse volume and warm peripheries

If no improvement

Repeat bolus of 20 ml/kg over 30 minutes, with careful monitoring for hepatomegaly, oxygen saturation, crepitation's in chest (if any of above appears then stop fluids)

If shock persists

Start Inj Adrenaline infusion @0.1 microgram/kg/min and refer to higher centre

#For severe acute malnutrition – consider SAM STW #For suspected Dengue follow Dengue Fever STW

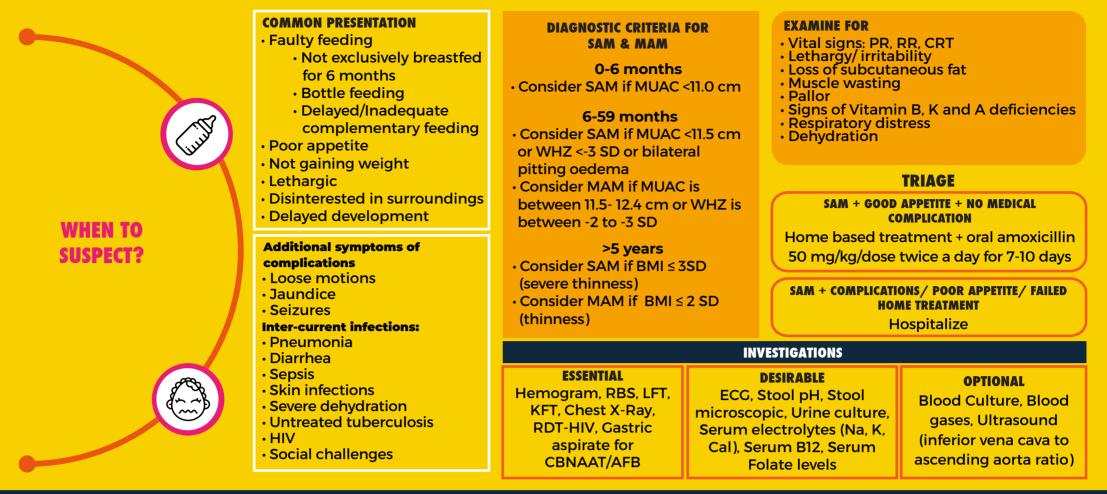
When to refer • Shock does not improve after 2nd fluid bolus • Signs of fluid overload • No facility for continuous monitoring. • Before referral counsel the parents and inform referring facility	When to Suspect Cardiac Failure • History of underlying heart disease • History of forehead sweating/ suck rest suck cycle • Murmur • Hepatomegaly or basilar crept If it is suspected be careful in giving fluid bolus		Complications Respiratory failure (excessive increase in the respiratory rates and inability to maintain saturation> 94% with oxygen) -non-invasive (CPAP/BIPAP) or invasive ventilation Congestive heart failure- Dobutamine / Milrinone infusion and Furosemide Infections on other sites- explore and treat accordingly 		
		DISCHARGE CRITERIA			
Completion of antibiotics as per culture sensitivity	Afebrile for 48 hours Vitals within normal limit for age		Good oral intake	Adequate urine output >1ml/kg/hr	
🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES					
		ing to Voice? If not; P Is the child respected at the child respected at the classify as danger similar to		s Unresponsive to voice (or being	







Standard Treatment Workflow (STW) for the Management of SEVERE ACUTE MALNUTRITION WITH COMPLICATIONS ICD-10-E43



TREATMENT

A. STABILISATION PHASE: Monitor vitals, urine frequency, stool/vomitus volumes INTAKE: IVF (DNS) 4 ml/kg/hr for 2-3 days with early/concomitant initiation of oral feeds (130 ml/kg/day)

INTARE: IVI (BNS)-	FIN/Kg/III IOI 2 5 days with a	early/conconnitant initiation of oral feeds (150 mi/k	(g/udy)			
CONDITION	PLACE OF TREATMENT	TREATME	NT			
INFECTIONS (empirically)	Facilities for supportive monitoring, investigations and IVF	 Inj Ampicillin - 50 mg/kg/iv or im X 6hrly Plus inj. Centamicin- 7.5 mg/kg iv or im, OD for 7-10 days If no response within 48 hrs or critically ill give inj. Ceftriaxone 50 mg/kg, OD for 7-10 days When accepting orally, switch to oral amoxicillin 40-45 mg/kg/dose twice a day for 7 days If prolonged diarrhea (>7 days): Metronidazole 10-12 mg/kg, 8 hrly for 7-10 days (inj.ectable or oral) 				
HYPOGLYCEMIA	Facilities for supportive monitoring, investigations and IVF	Conscious: 50 ml of 10% Dextrose or 1 tsf sugar in 3 tsf water orally				
(RBS <54mg/dL)	Transfer to intensive care facility to manage shock	Unconscious: 5 ml/kg of 10% Dextrose IV NO IMPROVEMENT treat as shock				
HYPOTHERMIA	Facilities for supportive monitoring, investigations and IVF. Plus warmer	Skin to skin care with mother (infants) Warming under warmer, incandescent lamp or warmer				
(<35.5 °C or 96 °F)	Intensive care facility to manage shock	NO IMPROVEMENT treat as shock				
Facilities for supportive monitoring, investigations SEVERE and IVF		Conscious: 50 ml of 10% Dextrose or 1 tsf sugar in 3 tsf water orally				
DEHYDRATION	Transfer to intensive care facility to manage shock	Unconscious: 5 ml/kg of 10% Dextrose IV NO IMPROVEMENT treat as shock				
ELECTROLYTE IMBALANCE (emperically)Facilities for supportive monitoring, investigations and IVF		Potassium: 3-4 mmol/kg/D, orally for 2 wks Magnesium: 0.4-0.6 mmol/kg/D1 IM followed by oral for 2 wks				
ANEMIA	Facilities for supportive monitoring, investigations and IVF	Whole blood /PRBC transfusion (10 ml/kg over 3 hrs): if H distress with close monitoring and hy. Furosemide (1 mg				
B. REHABILITATION PH	ASE (Transfer to NRC when cl	nild meets criteria for discharge [*] & accepts home av	ailable foods)			
	FEEDING ent: Facilities for supportive moni Treatment:	monitoring	VITAMINS Place of treatment: Nutritional rehabilitation center (NRC)			
increasing to give 150 switch to F100 for ne available food	F75 at least 5 times/day graduall 0-200 kCal/kg/day (usually 2-3 day xt 5-7days with introduction of ho ne as above with return to exclusi ever possible	b. Copper: 0.3 mg/kg/day X 2 wks orally	Treatment: a. Vitamin A: >12 months- 2 lac iu, 6-12 months: 1 lac iu, <6 months: 0.5 lac iu if food not fortified b. Vitamin D, A, B Complex: RDA			

*CRITERIA FOR DISCHARGE FROM HOSPITAL TO OUTPATIENT CARE: Clinically well and alert; no or resolving medical complications; no or resolving oedema (if present); satisfactory oral intake has a good appetite (taking at least 75% of target calorie intake of 150- 200 kcal/kg/day & 0-6 months old have weight gain of 3-5 gm/kg/day for three days).

PRIMARY FAILURE OF TREATMENT: (a.) Failure to regain appetite by day 4 (b.) Failure to lose oedema by day 4 (c.) Oedema still present Day 10 (d.) Failure to gain at least 5g/Kg/day for 3 consecutive days on catchup diet. Look for unrecognized congenital abnormality, inborn errors of metabolism, immune deficiency, other major organ dysfunction, and malignancy.

APPETITE TEST: Passed if, a child not fed for last 2 hours, when fed by mother in a quiet place consumes in 1 hour:

• 7-12 months: of \geq 25 ml/kg of F100

• > 12 months: of locally prepared ready to eat food **

AMOUNT TO BE GIVEN: 15 gms or more if < 4 kg; 25 gms or more if 4 – 7 kg; 35 gms or more if 7-10 kg **[Mixture of Roasted groundnut 1000 gm , Milk powder 1200 gms, Sugar 1120 gms, Coconut oil 600 gms. To be kept refrigerated for not more than 1 week.]

HOW TO PREPARE F75 AND F100	F75	F100
FRESH WHOLE CREAM MILK	300 ml	900 ml
SUGAR	100 gm	75 gm
VEGETABLE OIL	20 ml	20 ml
ADD WATER TO GET TOTAL VOLUME OF	1 Litre	1 Litre

ABBREVIATIONS

WHZ: Weight for Height Z-score **SAM:** Severe Acute Malnutrition

MUAC: Mid-upper Arm Circumference **SD:** Standard Deviation (from median)

MAM: Moderate Acute Malnutrition **BMI:** Body Mass Index

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

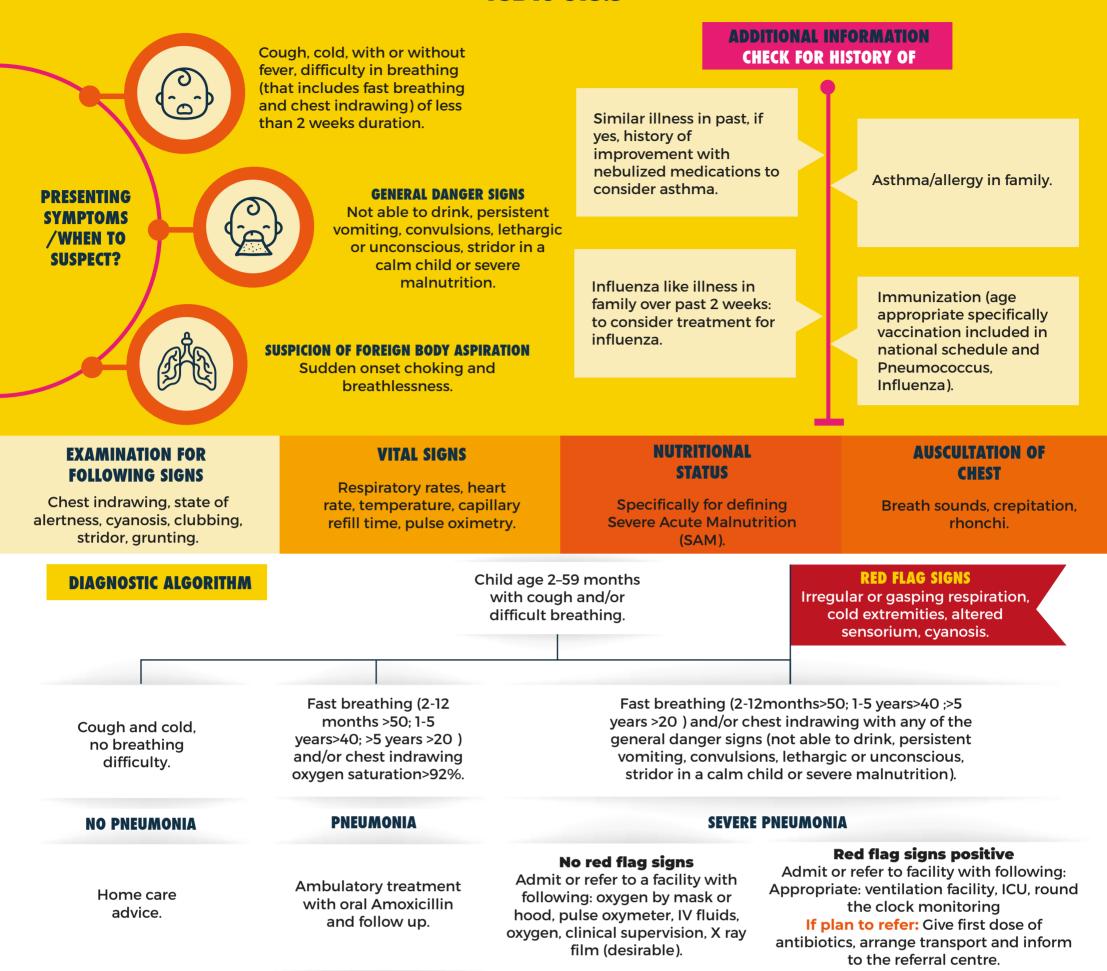
REFERENCES

- 1. The WHO growth standards. Available at http://www.who.int/childgrowth/standards/en/
- 2. Management of severe acute malnutrition in children 6-59 months of age with oedema. Available at http://www.who.int/elena/titles/oedema_sam/en/
- 3. Operational guidelines on Facility Based Management of Children with Severe Acute Malnutrition. Available at
- http://nhm.gov.in/nrhm-components/rmnch-a/child-health-immunization/child-health/guidelines.html
- 4. Kumar R, Kumar P, Aneja S, Kumar V, Rehan HS. Safety and Efficacy of Low-osmolarity ORS vs. Modified Rehydration Solution for Malnourished Children for Treatment of Children with Severe Acute Malnutrition and Diarrhea: A Randomized Controlled Trial. J Trop Pediatr. 2015 Dec;61(6):435-41.





Standard Treatment Workflow (STW) for the Management of SEVERE PNEUMONIA IN CHILDREN ICD10-J18.9



INVESTIGATIONS		TREATMENT			
ESENTIAL: Hemogram, random blood sugar, CRP, chest X-ray. DESIRABLE: Blood culture, pleural tap, serum electrolytes, renal and liver function tests. OPTIONAL: ABG, lung ultrasound, PCT, tracheal aspirate (gram stain with culture), bronchoscopy/BAL, microbiology culture, investigations for atypical organisms, PCR for viral etiology.	maintain oxygen satura IV ANTIBIOTICS: For children 2-59 months: Gentamicin ±5-7.5 mg/ For children >5 years: Am (Azythromycin/Erythrom) If suspected Staphyloco CXR, post measles, infe Amoxiclavulanic acid. SUPPORTIVE CARE: Parace (inhaled) as needed. WHEN AND WHAT TO SWITC Child is afebrile, RR has re indrawing and accepting of total of 5-7 days duratio If getting Doxacillin/Amor weeks. Start feeding as soon as	Ampicillin 100-200mg/kg in fc	 NON RESPONDERS: persistence of symptoms and/or signs 48-72 hours after initiation of appropriate treatment-change treatment-change antimicrobials. PLEURAL EFFUSION: diagnostic aspiration. PLEURAL EFFUSION: diagnostic aspiration. PMPYEMA: drainage with ICD. LUNG ABSCESS: change antibiotics for longer duration (4-6 weeks). PNEUMOTHORAX: Intercostal drainage. RESPIRATORY FALLURE: consider ventilation. INFECTION IN OTHER SITES: identify and treat appropriately. 		
ADDITIONAL	First and second line antibiotics for severe	FIRST LINE	ALTERNATE FIRST L	INE	SECOND LINE
INFORMATION	pneumonia:	Ampicillin	First gen Cephalosp	orins	Amoxiclav Cefuroxime Cefotaxime/ Ceftrioxone
WHEN TO REFER TO HIGHER CENTERS?	WHEN TO SUSPECT INFECTION WITH H1N1 VIRUS?	WHEN TO SUSPECT ACUTE BRONCHIOLITIS?	WHEN TO SUSPECT ASTHMA?		WHEN TO SUSPECT CHRONIC RESPIRATORY PROBLEM?
Facilities (as described above) for treatment or complications (if develops) are not available, suspecting chronic respiratory problems.	Child with cold, cough, fever with similar illness in any family members, consider H1N1 infection. Start Oseltamivir (as per national guideline).	A child below 2 years of age fulfilling case definition of first episode of severe pneumonia with predominant finding of wheezing on auscultation.	A child of age >3 years with history of recurrent cough, cold, wheezing with or without fever with good response to bronchodilator and personal or family history of asthma.		Child has any of the following: severe malnutrition, clubbing, feeding difficulty, family history of sibling death due to pneumonia, multi site infections (diarrhea, ear discharge oral thrush).

TDEATMENT

OMDUCATIONS AND THEID

Discharge when child is switched to oral medications, accepting oral for 24 to 48 hours

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

REFERENCES

- 1. Integrated Management of Childhood Illness (IMCI) (revised). Geneva, World Health Organization/The United Nation Children's Fund (UNICEF), 2014 (http://www.who.int/maternal_child_adolescent/documents/IMCI_chartbooklet/en/).
- 2. Revised WHO classification and treatment of childhood pneumonia at health facilities. http://apps.who.int/iris/bitstream/handle/10665/137319/9789241507813_eng.pdf;jsessionid=8BF6F1C94BD7BA81B8F464D4CBA40249?sequence=1
- 3. Bradley JS, Byington CL, Shah SS, et al. Executive summary: the management of community-acquired pneumonia in infants and children older than 3 months of age: clinical practice guidelines by the Pediatric Infectious Diseases Society and the Infectious Diseases Society of America. Clin Infect Dis 2011;53:617-30.
- 4. Lodha R, Kabra SK, Pandey RM. Antibiotics for community-acquired pneumonia in children. Cochrane Database Syst Rev. 2013 Jun 4;(6):CD004874



PSYCHATRY





Standard Treatment Workflow (STW) for the Management of ALCOHOL USE DISORDERS ICD10-F10

	s	pecial attention	to:	ASSESSA	NENT (DETAILED HISTORY)			
Universal screening for every patient attending any healthcare facility	(,	AUDIT can be used or screening) H/ o head injury Appearing under influence of alcoho H/ o impaired social, occupational functioning Daily alcohol consumption	 Age at initiation, quantity, frequency and progression (daily use and/or morning drivents) Age at initiation, quantity, frequency and progression (daily use and/or morning drivents) Time of last alcohol use and amount Binge drinking (men: 5 drinks over 2 hours; women: 4 drinks over 2 hours) Withdrawal state: insomnia, restlessness, anxiety, tremors. Use of alcohol (or benzodiazepines) to relieve or avoid withdrawal symptoms. Tolerance: increased doses of alcohol taken to achieve effects produced by earlier in Craving Difficulty in controlling duration of drinking or amount of use Preoccupation with alcohol use with neglect of alternative pleasures or interests Increased time spent to obtain/ take alcohol/ recover from its effects Continued use despite patient being aware of evidence of harmful consequences the occurred Abstinence and treatment attempts in past and reasons for relapse Co-morbid medical illness or psychiatric illness and their treatment Complications: Physical- gastritis, peripheral neuropathy, hepatic dysfunction, accidents/inju 					
		Drinking in large quantities			EXAMINATION			
		(men: 5 or more drinks/ day; women: 4 or more drinks/ day)	• BP • Pulse Rate	• Tremor • Sweating • Tachycardia	• Enlarged liver • Icterus • Abdominal swelling	 NEUROLOGICAL SIGNS Cerebellar signs Peripheral neuropathy Confusion 		
				DIAGNOSIS				
 Involvement in r binge drinking, o influence of alco It should have re 	Hazardous or Harmful useAlcohol dependence (three of the following six criteria to be present for at least one month)• Involvement in risky behaviours such as binge drinking, driving under the influence of alcohol1) A strong desire or sense of compulsion to take alcohol 2) Difficulty in controlling alcohol use 3) Withdrawal state when alcohol use has stopped or been reduced or use of the alcohol (or benzodiazepines) to relieve or avoid withdrawal symptoms• It should have resulted in harmful physical or psychosocial consequences4) Evidence of tolerance 5) Preoccupation with alcohol use 6) Alcohol use persisting despite clear evidence of harmful consequences							
				INVESTIGATIONS				
СВС	Liver fu	nction test	Blood sugar	Electrolytes	CT head (in case of se	zure/ delirium tremens)		
				MANAGEMENT				
Interventio • Alcohol De	azardous/ I on* to redu ependent u rate for trea	ARY CARE Harmful users - Br ice/stop consump users – Advice to s atment using Brie	tion top use		• H/ o withdrawal seizures/ hallucinations • Additional psychi	atric REFER TO SECONDARY		

SECONDARY CARE

- Treatment of withdrawal symptoms
- Managment of withdrawal seizure
 - Inpatient management with benzodiazepines (diazepam or lorazepam)
 - Frequent titration of medication. Higher dosage may be required.
 - Closer monitoring and nursing care

• H/ o delirium

Major medical

Additional substance

tremens

problems

use

- Treatment of additional psychiatric disorder or substance use disorder
- disorder
- Recurrent failed attempts at treatment

SECONDAR CARE IF

TERTIARY CARE

- Treatment of delirium tremens
 - R/ o head injury, hepatic encephalopathy, Wernicke's encephalopathy
 - \cdot R/ o other causes of delirium
 - Manage on similar lines as withdrawal seizures
 - Management in ICU setting when indicated
- Consult with other medical specialists (like gastroenterology or medicine for hematemesis).
- Management for suicidality or violence when emergent threat

***BRIEF INTERVENTION**

Inquire using open ended questions in a non-judgmental manner. Help patient to evaluate the risks versus the perceived benefits and to arrive at a decision to reduce or stop alcohol use. Includes (FRAMES) :

- Feedback about alcohol related problems
- Responsibility acknowledging that the patient is responsible for making the decision about their alcohol use
- Advice regarding the harms associated with continued use
- Menu of alternative change options (includes identifying alternative activities such as hobbies, involving the family in treatment)
- Empathetic attitude

REFER TO

TERTIARY

CARE IF

• Self efficacy - to encourage patients' confidence that they can make changes in their alcohol use and lifestyle

WITHDRAWAL MANAGEMENT

- Tab Diazepam (20-40mg/day in divided doses) based on severity of withdrawals.
- Monitor and titrate dose.
- If patient comfortable, reduce dose of medication by 10% to 20% per day, taper within 7 to 10 days
- Thiamine 100 mg OD
- Significant liver dysfunction: Lorazepam (2 mg Lorazepam equal to 5 mg Diazepam)

RELAPSE PREVENTION

(Long term goals- abstinence and socio-occupational integration)

- Disulfiram (250 mg OD) Pre-requisites:
 - Motivated patient
 - Patient's written consent
 - Under supervision of family members.
 - Inform patient and family about unpleasant, potentially serious reaction with even small amounts of alcohol (flushing, headache, vomiting, reduction of blood pressure, arrhythmias)
 - Ability of health personnel in the area to handle a potential reaction
- Relapse prevention counselling:
 - Identify cues leading to craving (like person, place, situation etc)
 - Develop strategies to deal with them effectively

INDICATIONS FOR ADMISSION

Failure of outpatient treatment

H/o withdrawal seizures/ delirium tremens

Co-morbid significant medical illness and/or psychiatric illness

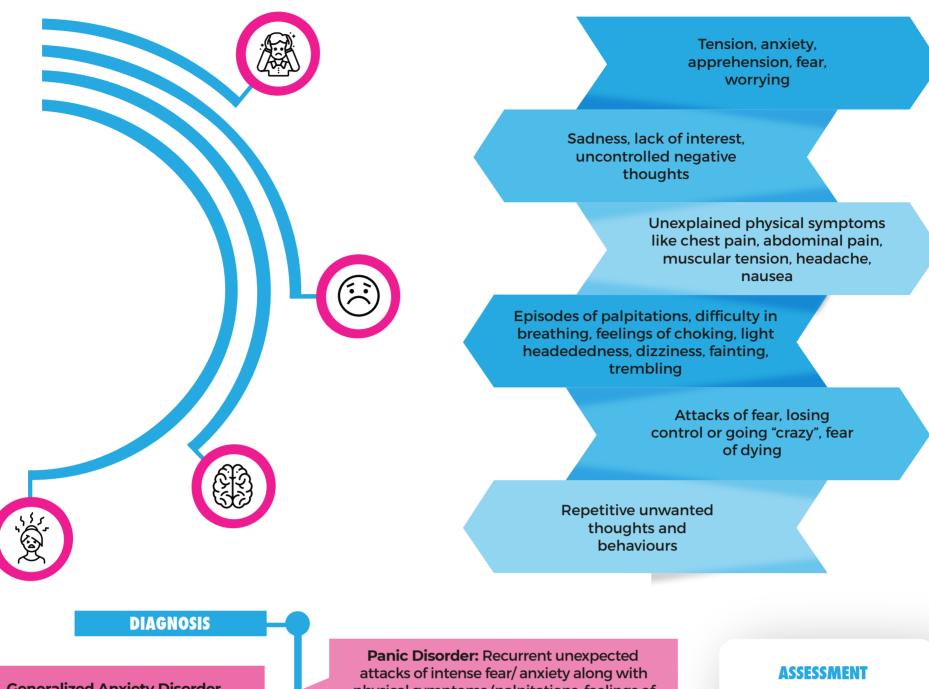
Poly-substance use

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



NDIAN COUNCIL O MENDICAL RESEARCI Serving the nation since 19

Standard Treatment Workflow (STW) for the Management of ANXIETY DISORDERS ICD-10-F40-F42



Duration of anxiety

- Degree of distress, and impairment of day-to-day functioning
- Symptoms of depression
- Substance and alcohol misuse
- Physical disorders: thyrotoxicosis, pheochromocytoma and hypoglycaemia
 Psychosocial factors: ongoing stress and other issues pertaining to work, family

Generalized Anxiety Disorder (GAD): Chronic feeling of tension, apprehension, anxiety or worrying about a number of events or activities that involve every day routine life circumstances (e.g., work, school, health, finance, household chores etc.)

Agoraphobia: Fear of going out of home alone, being in enclosed spaces (e.g., malls, cinemas etc.), open spaces (e.g., bridges, vast playgrounds etc.), using public transportation (e.g., trains, buses, planes etc.) Panic Disorder: Recurrent unexpected attacks of intense fear/ anxiety along with physical symptoms (palpitations, feelings of "choking", trembling, chest pain feeling dizzy/faint etc.)

Social Phobia: Marked fear and avoidance of social situations (e.g., interaction with strangers, meeting unfamiliar people, performing in front of others)

Obsessive-compulsive disorder (OCD): Recurrent and persistent unwanted thoughts (e.g., unwanted sexual and blasphemous thoughts, fear of harming self or others, fear of contamination, doubts about daily activities etc.) and repetitive behaviours (e.g., excessive washing / cleaning, checking, ordering etc.)

PRIMARY CARE LEVEL

Psychoeducation

- Reassurance
- Explain symptoms are of anxiety/ fear and mimic symptoms of physical illnesses (e.g., heart attack)
- Do not investigate excessively. Few investigations like ECG, ECHO maybe necessary in some patients
- Discourage doctor shopping
- Do not avoid triggers of panic attacks (e.g., physical exertion, agoraphobic situations) and fear (e.g., travelling by public transport).
- Emphasize avoidance maintains fears and phobias.
- OCD: Educate that the unwanted thoughts are a part of illness, and not a reflection of character or hidden intentions.

Pharmacological treatment

- Mild illness: Spending time, reassurance, and psychoeducation. May not need any medications.
- No improvement (few weeks): Escitalopram 5mg / day at night, with increase to 10 mg/d in a week. No satisfactory improvement in 4-6 weeks, may increase to 20 mg / day. If there is no significant improvement in another 4-6 weeks, refer to a specialist.
- Severe and unbearable anxiety: Diazepam (5 -10 mg) may be given at night. Do not continue for >1 month. Taper and stop over 2 weeks. Long-term treatment with benzodiazepines to be avoided
- Escitalopram to be continued for at least 1-2 years after remission
- Side-effects (sexual dysfunction, sedation, weight gain): monitor and address periodically

MANAGEMENT

SECONDARY CARE LEVEL (DISTRICT HOSPITAL)

- Review diagnosis and treatment history if there is no improvement with a trial of Escitalopram.
- Check whether the patient has taken medication at prescribed dose and on a regular basis
- Second SSRI
- (either of them for about 2-3 months):
 - Sertraline upto 200 mg/day,
 - Fluoxetine upto 60 mg/day,
 - Paroxetine upto 50 mg/day,
 - Fluvoxamine upto 300 mg/day
- No response to second SSRI: cognitive behaviour therapy (CBT) if trained therapists available.
- Refer to tertiary centre if unsatisfactory response after second SSRI and / or addition of CBT.
- If referral to tertiary centre is not feasible, psychiatrists may try other strategies (other than Deep Brain Stimulation and surgery for OCD) mentioned under the "tertiary care" at the secondary level itself.

TERTIARY CENTRE (MEDICAL COLLEGE, REGIONAL MEDICAL CENTRE, PSYCHIATRIC HOSPITAL)

· Evaluate reasons for treatment resistance like

- Wrong diagnosis
- Inadequate drug treatment,
- Poor adherence to treatment
- Inadequate CBT,
- Presence of comorbid conditions such as personality disorders and organicity
- Panic disorder: evaluate any medical conditions that mimic panic disorder (hyperthyroidism, hyperparathyroidism, pheochromocytoma, vestibular diseases, seizures, arrhythmias, etc.).
- OCD: Trial of third SSRI or clomipramine
- Treatment resistant OCD: inpatient treatment for intensive therapist-assisted daily CBT and for rationalization of medication regimen.
- Other anxiety disorders: Trial of non-SSRIs (e.g., venlafaxine, duloxetine, pregabalin etc.) and tricyclic antidepressants
- If response to medications is poor or unsatisfactory:
 - CBT is the preferred mode of treatment alone or in combination with medications.
 - Treat comorbid psychiatric disorders (e.g., personality disorders)
 - Pharmacological augmenting strategies if antidepressants and CBT do not provide relief.

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES







Standard Treatment Workflow (STW) for the Management of **CHILDHOOD BEHAVIORAL DISORDERS** ICD10- F90-98

OPPOSITIONAL DEFIANT DISORDER (ODD)

 Doesn't obey or listen, back-answers, rude behaviors Demanding, stubborn, throws tantrums when demands are not met

CONDUCT DISORDER

 Aggressive – angry, abusive, fights, hits or hurts people, bullies other children, damages articles • Stealing, lying, threatening or misbehaving with people, truant (keeps away from school without parents' knowledge), runs away from home, bad company, cruelty to animals

ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)

- Restless, always on the move / running, can't sit in a place,talkative
- Can't focus attention on a task, poor concentration, gets easily distracted, disorganized, does not complete school work
- Too mischievous, can't be left alone, troubles people or damages things, or gets injured Impatient, always in a hurry, can't wait for his turn, does not care for danger, acts without thinking

ALL 3 DISORDERS

Poor or erratic school performance and / or complaints about behaviour from school

DIAGNOSIS

Symptoms are present and persistent over several months Attempt further classification into ADHD, ODD and CD (ADHD may be present with or without ODD and CD

CAUTION

- Abrupt onset
- Recent onset (few weeks to few months)
- Sudden increase in severity (consider another psychiatric disorder such as bipolar affective disorder, hypomanic or manic episode -> follow the relevant STW)

PRESENTATION

CLINICAL

PARENT INTERVIEW

- Symptoms- onset, duration, type(ODD, Conduct, ADHD-as above) and severity
- Developmental problems, emotional disturbances and stress
- Alcohol and substance use /misuse
- Impact on child and family

PSYCHOEDUCATION

ASSESSMENT (History From Multiple Sources)

- **FAMILY SITUATION**
- Health (including mental health) and wellbeing of family members
- Cohesion, mutual understanding and harmony in the family
- Parenting and childrearing practices: caring and disciplining, criticism, unfair comparison and physical punishments, mutual blaming of parents for child's problem

SCHOOLING Attendance

- Performance
- Learning
- problems, Classroom
- behaviors Recent changes
- in syllabus and/or school

CHILD INTERVIEW Develop rapport(discuss neutral topics; avoid direct

- tackling of misbehaviors)
- Observe:
 - Features of ADHD (restless, fidgety, easily distracted, attention keeps shifting)
 - Speech and language ability, intelligence, academic skills and mood
 - · Enquire about any stress or difficulties child is facing at home, school, and with peers and anger control

MANAGEMENT

- Avoid advice
 - Anger management (count from 10 -1 backwards, move away from situation, deep breaths, relax, self-talk to cool down)

WORK WITH THE CHILD

Children with ADHD:

WORK WITH THE SCHOOL

- Feedback to school regarding child's
- Teachers to give extra attention, help and support for the child
- Extra coaching, if

- ventilate, validate and empathize their difficulties, reassure)
- Recognize and manage mental health problems such as depression and alcohol

Help parents deal with their own worries and stress (listening, giving space to

WORK WITH FAMILY

- Multifactorial causes-lack of self-regulation, and adverse environment

- Parents can directly contribute to the child's improvement

- Explain the child's behaviours are not intentional

- Not child's fault, do not blame the child

- Can be improved with proper management

- - condition

problem in parents

Parent management training*

*PARENT MANAGEMENT TRAINING

- Analyse the problem behaviors and understand patterns : time of occurance, triggers, duration and consequencies
- Engage with child in mutually enjoyable, pleasurable activities (playing games, discussing interesting things or doing activities together)
- Set clear do's and don'ts and explain to child in clear, simple, short instructions the consequencies (like withholding privileges following misbehavior; use star-charting (contingency management) and rewards based on number of stars earned
- In children with ADHD, develop clear daily routines, supervise activities and appreciate on completion of taks
- Limit screen time/ monitor use of electronic devices

• Dos

- Consistency in enforcing rules
- Catch the child being good and praise
- Ignore negative behaviours
- Child can be put in a boring place till he/ she becomes quiet for a few minutes (time-out)
- Encourage age appropriate responsibilities
- Don'ts
- Bribe
- False promises and threats
- Harsh punishments
- Excessive criticism and blaming especially in front of others
- Unfair comparison
- Yielding to unreasonable demands

"stop-think-act" or "halt and proceed" technique

needed in case of learning problems

MEDICATION (AVOID BEFORE 5 YEARS)

Severe and persistant aggression:

- **T. Risperidone** under close supervision (starting dose-0.25 mg, single daily morning dose after breakfast. Based on response, increase by 0.25 mg weekly up to 1 mg single daily dose).

- Not to exceed 1 mg/day
- **Response + :** continue 3 months f/b slow taper
- Response -: 4 weeks trial, then refer
- Monitor adverse effects: weight gain, extra-pyramidal symptoms (EPS)
- [if EPS : add I mg Trihexyphenidyl OD morning]
- Severe hyperactivity and impulsivety:
 - T. Clonidine (starting dose-25 µg single daily dose before sleep, increase by 25 µg weekly up to100 µg per day in 2-3 divided doses
- Monitor BP and drowsiness
- Advise against sudden discontinuation

REASONS FOR REFERRAL

Severe aggression Severe, complicated presentation Lack of response to treatment Highly dysfunctional family Alcohol and substance abuse

SECONDARY CARE (DISTRICT HOSPITAL)

- Review and reassess diagnosis (clinical evaluation using Rutter's multi-axial system) and all the pointers given above
- If failed trial of Clonidine/ Moderate ADHD: T. Atomoxetine (starting dose-10 mg single daily morning dose after breakfast. Increase up to 1mg/kg/day under close supervision).
- Monitor adverse effects and response
- Systematic parent management training / behavioral management and individual therapy (as given above)
- TERTIARY CARE (MEDICAL COLLEGE / REGIONAL REFERRAL CENTRE)
- Evaluate and manage severe behavior disorders severe ADHD, ODD, and CD, if necessary on short-term inpatient basis
 - Multi-modal management with clear individualized plan
 - Trial of Methylphenidate in moderate / severe ADHD under expert supervision
 - · Recognize and treat comorbid disorders such as bipolar disorder, substance use disorder, and internalizing disorders and manage
 - Pharmacological management of older children / adolescents with severe aggression / impulsivity with Risperidone and/or Lithium
 - Family therapy for dysfunctional / discordant families, contributing to child's condition
 - Management of children in difficult circumstances with mental health issues (children in need of care and protection; children in conflict with law)

REFERENCES

• World Health Organization. mhGAP intervention Guide-Version 2.0 for mental, neurological and substance user disorders in non-specialized health settings. Geneva: WHO. 2016.

Pliszka S, AACAP Work Group on Quality Issues. Practice parameter for the assessment and treatment of children and adolescents with attention-deficit/hyperactivity disorder. Journal of the American Academy of Child & Adolescent Psychiatry. 2007 Jul 1;46(7):894-921.

Steiner H, Remsing L. Practice parameter for the assessment and treatment of children and adolescents with oppositional defiant disorder. Journal of the American Academy of Child & Adolescent Psychiatry. 2007 Jan 1;46(1):126-41.

🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES





Standard Treatment Workflow (STW) for the Management of CHILDHOOD EMOTIONAL DISORDERS

CLINICAL PRESENTATION -Recent Onset Behavioral Changes

SOMATIC (PHYSICALLY UNEXPLAINED) SYMPTOMS

- Weakness and tiredness
- Aches and pains
- Headache
- Non-epileptic attacks of fainting
- Chest pain and stomach pain
- Hyperventilation- often triggered by stress or distress

SYMPTOMS OF DEPRESSION

- Loss of interest in usual activities
- Recent deterioration in school performance
- Wanting to be alone, withdrawn, not interacting with people
- Looks unhappy, "off mood", crying for trivial or no reason, irritable, sensitive to any criticism
- Decreased sleep, loss of appetite and weight loss
 Talking about death and dying, self harm (eg.
- self-cutting) or suicidal attempt

SYMPTOMS OF ANXIETY

- Always worrying, tense
- Exam tension, performance anxiety, worries about marks and ranks
- Excessive fear and avoidance of some objects or situations (insects, animals, ghosts)
- \cdot Reluctance or refusal to go to school
- Very shy, avoids social situations, scared of talking or interacting with strangers,
- Clinging to mother, scared of being separated from mother

DIAGNOSIS

- Persistent symptoms of emotional disturbance for several weeks, significantly affecting the child's life
- Unexplained by medical condition such as hypothyroidism
- Depression and anxiety symptoms can co-occur
- Depression more common in adolescents, may have features similar to adult onset depression

CAUTION

Assessment of suicidal risk and a plan of action is important in children with emotional disorders, especially depression (refer to appropriate STW) Elicit h/o hypomania/mania in children with moderate to severe depression (consider diagnosis of bipolar disorder) Physical conditions can cause similar symptoms (anemia and thyroid disturbance)

PARENT INTERVIEW AND HISTORY TAKING

- Onset, duration, severity and full range of symptoms
- Home environment, family life and relationships, parenting practices and stressors
- Information (from paretns and school) about school performance, behavior, school refusal, bullying experiences, peer relations and any recent change

ASSESSMENT

CHILD INTERVIEW

- Develop rapport
- Ask subjective distress (low mood, irritability, sadness, lack of enjoyment of activities, worries, fears, tensions, autonomic symptoms)
- Stressful events (loss, death in the family, separation, frightening experiences, traumatic abusive or shocking events, humiliating experiences, bullying in school, academic stress) and interpersonal difficulties
- Explore parent-child relations and interactions and any undue punishment or criticism

MANAGEMENT

WORK WITH THE CHILD

 Psycho-education of the child- explain they are suffering from an emotional problem and it is not their fault and they will get better with proper treatment

PHYSICAL EXAMINATION

(Rule out)

- Post-viral syndrome
- Recurrent attacks of
- malaria
- Chronic infections, chronic physical illness, anaemia, PCOD or thyroid disturbance

WORK WITH SCHOOL

• Give feedback to the school about

 Child is emotionally disturbed and not able to function well

WORK WITH PARENTS

- Not the child's fault

PSYCHOEDUCATION:

- Avoid undue criticism, over expectation, unfair comparison, scolding and punishment
- Parents' support, encouragement and understanding is important
- Counsel about suicidal risk in depression and to be alert to pointers to suicidality
- Evaluation and management of the mental health issues in parents
- Discuss about specific steps to reduce undue stress
 the child is facing
- Anxiety management and emotional regulation skills
 - Muscle relaxation
- Deep breathing exercises
- Praanaayaama / yoga
- Substituting distressing thoughts with more comforting thoughts
- Counsel the child to confide any distressing thoughts, including thoughts of death and dying
- Encourage the child to gradually return to the usual life and activities in a step-by step manner with parental support and encouragement
- child's condition and stress, need for support, encouragement and school's cooperation.
- If school refusal, graded return to school: encourage child to return to school gradually with the support of family and cooperation of school (e.g. initially for a few minutes in school compound, later for 1 period in school and moving on to longer duration

MEDICATION (MODERATE CASE OF DEPRESSION OR ANXIETY IN ADOLESCENTS)

- Tab Fluoxetine start at 10 mg OD morning, increase to 20 mg OD after 2 weeks depending on response
- Inform adverse effects: behavioral activation (marked restlessness and irritability), onset of hypomanic symptoms, and worsening of suicidal ideas. Stop drug if they are troublesome
- Avoid benzodiazepines (except as temporary measure for few weeks in severe anxiety attacks or panic attacks Clonazepam 0.25-1 mg /day)

SECONDARY CARE (DISTRICT HOSPITAL)

- Review and reassess diagnosis through detailed clinical examination using Rutter's multi-axial system
- Review the treatment received and plan multi-modal treatment.
- Reconsider medications, and augmentation strategies
- $\boldsymbol{\cdot}$ Review child's and family's awareness of the illness and do psycho-education
- Ascertain the presence of psychosocial factors : disturbed home environment, parent-child relationships and severe stressors
- $\boldsymbol{\cdot}$ Screen parents for mental health problems and manage accordingly
- **Individual therapy** focussing on identifying and challenging negative thoughts, anxiety management and coping with stress, helping them face difficult situations in small steps, improving interpersonal relationships
- **Parent counselling** to address family issues, communication and interaction patterns
- Collaborate with school wherever necessary (get school report; explain problem in simple terms, and suggest ways by which school can help)
- Recognize and manage less common problems such as obsessive compulsive disorder, psychoses and bipolar disorders
- Manage adolescents with mild / moderate suicidal risk

REASONS FOR REFERRAL

- Frequent expression of suicidal ideation/ attempted suicide / self-harm behavior such as self-cutting
- Severe symptoms
- Complicated picture, or features of obsessive compulsive disorder (OCD)
- No response to interventions in 4-6 weeks

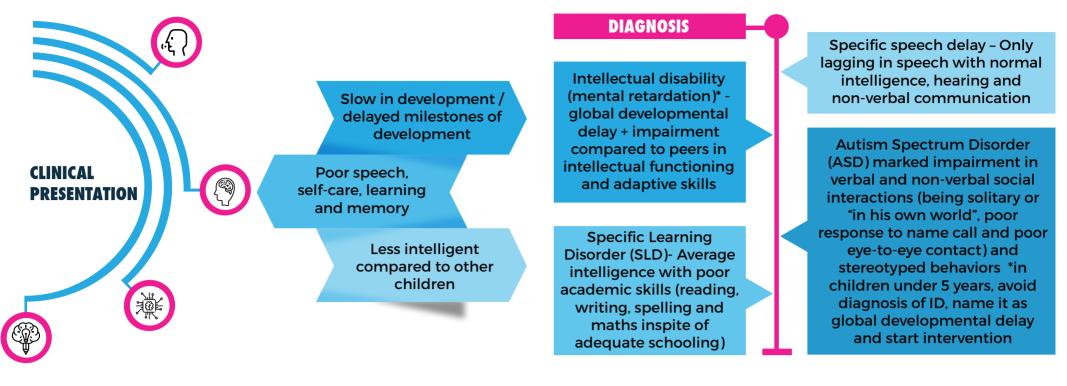
TERTIARY CARE (MEDICAL COLLEGE / REGIONAL REFERRAL CENTRE)

- Thorough diagnostic evaluation
- Manage severe mental disorders psychoses, recurrent mood disorders, adolescents with severe depression, & treatment resistant cases, persistent suicidality, recurrent self-cutting, if necessary in inpatient setting
- Family therapy for dysfunctional / discordant families contributing to child's condition
- Cognitive behavior therapy for older children with severe OCD, depression, and anxiety disorders
- ECT on case to case basis (older adolescents with severe depression, mania, psychosis or catatonia unresponsive to adequate pharmacological management)
- Appropriate psycho-social steps if there is abuse, maltreatment or neglect
- Neurology referral in suspected cases of epilepsy and organicity

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



Standard Treatment Workflow (STW) for the Management of **CHILDREN WITH DEVELOPMENTAL PROBLEMS** ICD10-F70-89



ASSESSMENT

DETAILED DEVELOPMENTAL PHYSICAL EXAMINATION: BEHAVIOURAL PROBLEMS: OTHERS: ASSESSMENT: • Height and weight, Hyperactive Family situation Head circumference. Assess if child is lagging behind Impulsive behaviors · Parents' awareness of Vision and hearing Sleeping and feeding problems in developmental attainments the child's problems compared to same-age children Any noticeable physical anomalies Aggression Quality of attention (club-foot) or unusual facial appearance Ask mother to estimate the and care being given Motor abnormalities (stiffness / spasticity or **EMOTIONAL PROBLEMS:** mental age of child to the child, weakness of limbs, unsteady gait) Ascertain if delay is global (all Excessive crying Past consultations and · Any other problems (heart murmurs, milestones) or restricted to one Irritability treatment area (motor or speech) organomegaly) Shyness and fears educational history

PSYCHO-EDUCATION OF PARENTS

- Normal reassure parents
- Mild delay ("at risk") early intervention and follow-up
- Explain causation due to some damage to brain before, during or after birth
- No medication can improve intelligence
- Teaching and training to improve skills and gaining independence
- Systematic, persistent and repetitive training as per the child's ability
- Treatment of associated problems (vitamin or mineral deficiency or

MANAGEMENT

EARLY INTERVENTION / SENSORY-MOTOR STIMULATION FOR **YOUNG CHILDREN – UNDER 3 YEARS**

- Create opportunities for the child to learn with interset and attention
- Engaging and spending time with child in activities
- Offer appreciation
- Engage the child to use eyes and ears (different types of sounds and sights), touch (eg., tickling, stroking, gentle massaging), movements (gentle movement of limbs, gentle bouncing, range of movement exercises) and improving hand functions (taking, holding, giving, pushing, pulling)
- Use play materials-rattles, paper balls, rubber balls, clay, soft dough, water play, soap bubbles, vegetables.
- Parallel vocalization to improve utterances (making

HOME-BASED PARENT MEDIATED SKILLS TRAINING

- Develop and maintain regular, stimulating daily routines
- Teach parent to teach child : simple imitation, pointing, pretend-play; self-help skills (eating, toilet training, bathing, dressing), doing simple household chores (washing utensils, helping in cleaning house), social skills - skills of interaction, simple academic skills, simple vocational skills, helping in kitchen under supervision, self-protection
- Find current level of adaptive abilities of the child and choose a target skill
- Tell and show how to do things (modelling), make the tasks simpler, break activities in simple

epilepsy, ADHD, vision/hearing issues,) - refer to appropriate STW

 Avoid overprotection, overindulgence and understimulation

EDUCATION AND TRAINING

- · Liaise with schools and ensure child attends school that is most appropriate
- Assist in enrolment to special school
- Consider training in vocational skills (informal and formal) for older adolescents

the same sound as the child immediately).

 Improve conceptual skills by classifying, arranging, sorting, and recognizing and naming activities (for eg., vegetable sorting, grain sorting, arranging vessels by their size and shape)

steps and teach one step at a time, notice and praise even minor efforts and improvements (rewarding or reinforcing), using hand-on-hand techniques (keeping your hand on the child's hand and making them do the activity)

SOCIAL WELFARE / LIAISON MEASURES

- · IQ testing and certification for social welfare benefits
- Help parents to link with other agencies/ services that deal with such children such as CBR programs or parent associations

- Severe or multiple developmental problems
- History of regression (loss of acquired skills)
- Definite family history of developmental problems (h/o similar problem in the sibling)

- Co-occuring severe behavioral or emotional problems
- Suspected case of ASD
- Suspected SLD
- Genetic counselling
- Speech therapy or physiotherapy

SECONDARY CARE (DISTRICT HOSPITAL)

- Psychological testing for ID, SLD and diagnosis of ASD
- Basic management of ASD home-based parent-mediated training in social, communicative, and self-help skills
- Appropriate management of behavior problems with medication / psychosocial or behavioral intervention (see relevant STW's)
- Help parents access relevant services such as District Early intervention centres (DEIC's), parent organizations, and benefits

TERTIARY CARE (MEDICAL COLLEGE / REGIONAL REFERRAL CENTRE)

- Evaluate and manage children with severe IDD, ASD, multiple disabilities, and those with severe comorbid disorders such as ADHD, aggression, bipolar disorder, and psychotic disorders through multi-disciplinary approach
- Investigate for the cause review tests already done; imaging, genetic tests, metabolic tests (as per requirement); arrange for genetic counselling
- Manage treatable disorders (like hypothyroidism and inherited metabolic disorders)
- Manage comorbid physical health problems (like epilepsy, visual /hearing impairment, locomotor/ orthopaedic problems)
- Assessment and management for SLD psychoeducation of the child and parents, liaison with school, teaching basic remediation techniques to parents, helping parents access relevant organizations, issue of exemption certificates, and decisions about further schooling such as open schooling

REFERENCES

- World Health Organization. mhGAP intervention Guide–Version 2.0 for mental, neurological and substance user disorders in non-specialized health settings. Geneva: WHO. 2016.
- Szymanski L, King BH. Practice parameters for the assessment and treatment of children, adolescents, and adults with mental retardation and comorbid mental disorders. Journal of the American Academy of Child & Adolescent Psychiatry. 1999 Dec 1;38(12):5S-31S.
- Girimaji SC.(2008) Clinical Practice Guidelines for the Diagnosis and Management of Children With Mental Retardation. Retrieved from www.indianjpsychiatry.org/cpg/cpg2008/CPG-CAP_05.pdf

🖝 KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

TThis STW has been prepared by national experts of India with feasibility considerations for various levels of healthcare system in the country. These broad guidelines are advisory, and are based on expert opinions and available scientific evidence. There may be variations in the management of an individual patient based on his/her specific condition, as decided by the treating physician. There will be no indemnity for direct or indirect consequences. Kindly visit our web portal (stw.icmr.org.in) for more information. © Indian Council of Medical Research and Department of Health Research, Ministry of Health & Family Welfare, Government of India.

REASONS FOR REFERRAL





Standard Treatment Workflow (STW) for the Management of DEPRESSION ICD10-F45

CORE SYMPTOMS

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CLINICAL DIAGNOSIS OF DEPRESSION	Le contraction de la contracti

Depressed Loss of Easy 2 mood fatigability/ interest diminished and activitv enjoyment **ADDITIONAL SYMPTOMS** Reduced concentration and attention Reduced self-esteem and self-confidence Ideas of guilt and unworthiness Bleak and pessimistic views of the future · Ideas or acts of self-harm or suicide Disturbed sleep Diminished appetite

To make a diagnosis of depression, symptoms must present for at least 2 weeks.

Severity of depression	Core symptoms	Additional symptoms
Mild depression	2	2 or more
Moderate depression	2	3 or more
Severe depression	3	4 or more

Rule out Bipolar Disorder / Grief / Adjustment Disorder

AT PRIMARY CARE

MILD DEPRESSION

- Advise Behavioral Activation to patients
- Practicing activity monitoring write down your activities / rate your depression / schedule activities that make you feel good / make a to do list/ set clear and specific goals
- Focusing on your value categories make time for your family / friends / set clear goals at work / contribute
 to community
- Reccomend yoga & mediatation
- Handling daily task monitor sleep /diet and practice good personal hygiene
- Supportive psychotherapy / Brief Counselling
- Validate the problems and ensure frequent follow-up
- If no improvement in 4 to 6 weeks, consider pharmacotherapy

REFERRAL TO SECONDARY CARE

BROAD MANAGEMENT PLANS

CLINICAL ASSESSMENT

Cognition

- Hopelessness (about future)
- Helplessness (about others)
- Worthlessness (about self)

Assessment of Suicide Risk

- Suicidal thoughts
- Suicidal idea
- Suicidal intent
- Immediate risk for attempt

INVESTIGATION

Assessment

of

Depressive

Cognition

Assess

friend

and

family

support

- Haemogram
- Thyroid function tests
- Electro Cardiogram
- Electrolytes (Sodium)
- Rule out secondary medical cause of depression like Hypothyroidism
- Rule out use of anticancer drugs
- (Cyclophoshamide) / anti retroviral drugs (Efavirenz, Zidovudine)/ Antibiotics (Dapsone, Ethambutol)/ Anabolic Steroids/ Propanolol • Rule out associated comorbid medical condition – Diabetes, Stroke, Epilepsy, Cancer, Coronary Artery Disease and Auto Immune disorder

MODERATE / SEVERE DEPRESSION

- Tab Escitalopram 10 mg-20 mg /day or Cap. Fluoxetine 20mg -40mg /day
- Tab. Clonazepam 0.25mg 0.5mg /day for sleep disturbance / anxiety symptoms and consider taper and stop after 2 weeks.
- If patient responds to SSRI in 2 to 4 weeks, then continue treatment for 6 to 9 months and taper and stop

AT SECONDARY CARE

- Difficulty in making diagnosis
- No improvement after 4 to 6 weeks of treatment with first line medications
- Depression in special population: Elderly / Pregnancy / Lactation / Children / Adolescents
- Comorbid medical illness /
 Substance use
- Suicidal risk assessment

REFERRAL TO TERTIARY CARE

- No improvement in 2nd line treatment
- Immediate risk for suicidal attempt / thought
- Needing intense counselling/ psychotherapy
- Co Morbid Substance -Cannabis / Poly substance

- Selective Serotonin Reuptake Inhibiters (SSRI) are usually first choice (watch for GI bleed and drug interaction)
- Improvement starts in in 2nd week and expect
 adequate response by 6 weeks
- Duration of treatment typically lasts 6-9 months and Gradual tapering of medication advised for first episode
- Restart SSRI , In case of resurgence and recurrence
 of depressive symptoms
- Observe for switch / activation with Antidepressants
- Watch for risk of overdose with TCA (Amitriptyline / Imipramine) and Mirtazapine

SPECIAL POPULATION

- Pregnancy / Lactation period -Pre Conception counselling and preferred drug is
 Tab. Sertraline 50 mg - use lowest possible dose
- Elderly -
- Tab. Escitalopram 10 -20 mg or Tab. Sertraline 100 mg
- (monitor for hyponatremia)
- Avoid TCAs like Amitriptyline / Imipramine in Elderly (due to anticholinergic side effects)
- Adolescents- Cap. Fluoxetine
- 20 -40 mg /day (observe for switch / activation/ suicidality)

- Confirm Diagnosis and Suicide risk
 assessment
- Assess for other Medical
 Comorbidities
- Investigations Haemoglobin, Thyroid Function Test, Electrocardiogram
- Non Responder Switch over to SNRI (Venlafaxine 75 – 150 mg, Mirtazapine 30 mg) or TCA (Amitriptyline 75 -225mg / Imipramine 75 -225mg)
- Cognitive Behavioral Therapy /
 Problem Solving Therapy
- Add on Yoga Therapy / Meditation

AT TERTIARY CARE

- Reconfirm Diagnosis
- Assess other psychiatric comorbidities
- Partial Responder Optimise the SNRI /TCA or Augment with Tab. Lithium 300 to 600mg /per day or Tab. Thyroxine 25 - 50 ug per day.
- Non Responder Add Tab. Sertraline 100mg or Tab. Bupropion 300mg to existing Venlafaxine 150mg / Tab. Mirtazapine 30mg / Amitriptyline 225mg / Imipramine 225mg.
- Add on Electro Convulsive Therapy for Catatonia / Suicidality
- Add on Cognitive Behavioural Therapy/ Inter Personal Therapy / Problem Solving Therapy
- Add on low dose antipsychotic treatment (Risperidone 2 -4 mg / Tab. Olanzapine 5 - 10 mg) for psychotic symptoms

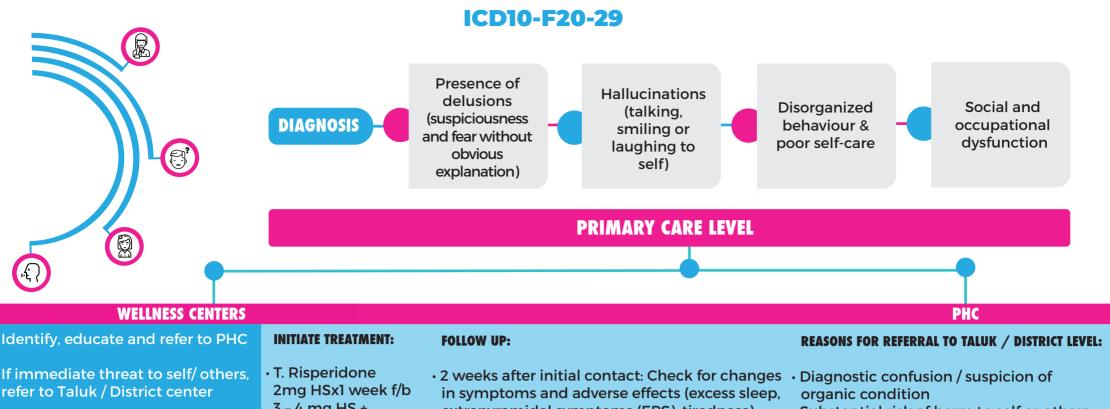
REFERENCES

- Gautam S et al., Clinical Practice Guidelines for the management of Depression. Indian J Psychiatry. 2017;59(Suppl 1):S34-S50.
- Avasthi A, Grover S. Clinical practice guidelines for management of depression in elderly. Indian J Psychiatry 2018;60, Suppl S3:341-62
- Sarkar S, Grover S. A systematic review and meta-analysis of trials of antidepressants in India for treatment of depression. Indian J Psychiatry. 2014;56:29–38
- National Institute for Clinical Excellence. Depression: management of depression in primary and secondary care. Clinical Guideline 23. London: NICE, 2004.
- mhGAP Intervention Guide Version 2.0 for mental, neurological and substance use disorders in non-specialized health settings. World Health Organisation, 2016

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



Standard Treatment Workflow (STW) for the Management of **PSYCHOSIS**



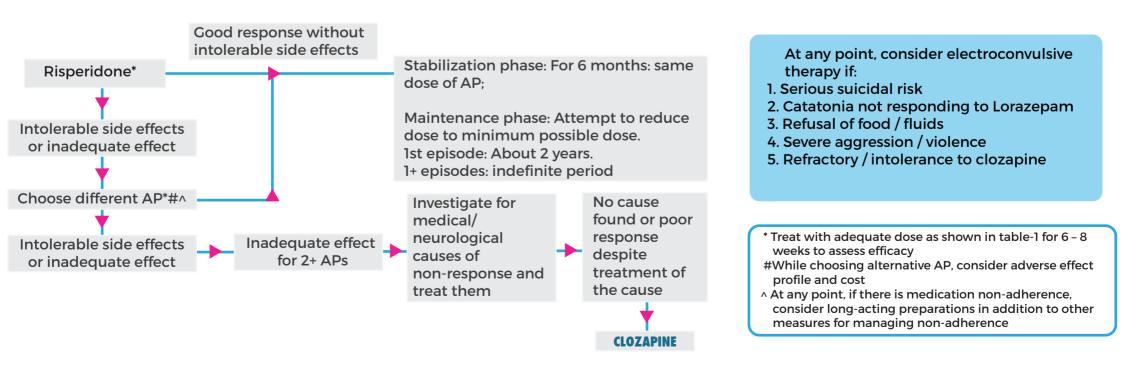
FOLLOW-UP AND REHABILITATION:

- Monitor & manage challenges in treatment continuation If unsatisfactory outcome despite regular treatment:
- Liaise with higher centers for optimal outcome
- Liaise with social welfare department for disability certification & welfare benefits if continued poor outcomes
- 3 4 mg HS + Trihexyphenidyl
- (THP) 2mg(morning) • Psychoeducation: - medical model of
- psychosis - address misconceptions &
- build hope - inform about possible adverse effects of medications
- extrapyramidal symptoms (EPS), tiredness) adjust the dose of risperidone and THP accordingly; address questions if any; advise gradual return to work/school; give specific follow-up date; liaise with wellness center for ensuring continuity of care
- Once in 1 2 months: Check for symptoms, functioning and adverse effects (EPS. weight-gain, menstrual/sexual dysfunction); adjust the dose of Risperidone (range: 2 - 8 mg/day) and THP (range 2 - 6 mg/day); liaise with wellness center for ensuring continuity of care
- Substantial risk of harm to self or others and catatonic symptoms
- · Comorbid substance use, depression/anxiety, intellectual disability
- · Poor symptom-control or functioning despite regular treatment or poor treatment adherence
- · Significant adverse effects: weight-gain, metabolic adverse effects, tardive dyskinesia
- Questions regarding marriage, pregnancy, sexual dysfunction

SECONDARY CARE (TALUK/DISTRICT HOSPITALS)

INDICATION FOR REFERRAL FROM PHC	Diagnostic confusion	Poor response to Risperidone	Intolerance to Risperidone	adherence to	Comorbid conditions	Challenging situations	Rehabilitation needs	Pregnancy	
MANAGEMENT #Encourage follow up in primary care after addressing referral issues * Watch for adverse effects as SSRIs may increase serum levels of antipsychotics		Positive symptoms: Follow algorithm Negative symptoms: Rule out or manage depression/anxiety and extrapyramidal symptoms; Family counseling if understimulated/ over-protected Consider less-sedating antipsychotics and adding SSRIs*	Follow algorithm	 Assessment of factors causing poor adherence & specific manage- ment Consider depot anti- psychotics Liaise with primary care for assertive follow up 	 Depression/ anxiety: Brief psychological intervention; consider SSRIs* Substance use: Detoxification and brief interventions (see SUD module) Developmenta I disabilities: Behavioral 	 Suicidality: Inpatient care, Crisis management, Management of comorbidity; Consider ECT Violence: Verbal de-escalation IV sedation, Brief inpatient care 	 Assess disability & counsel about welfare benefits Rehabilitation counseling Family intervention for expressed emotions and attitudes & behaviors interfering with functioning Brief interventions for cognitive & social-skill deficits Address vocational/ educational challenges involving governmental/ non-governmental 	 Proactively address sexual and endocrine problems when relevant Educate about risk of obstetric outcomes, risk of relapse & risk of psychosis in the offspring 	
				TERTIARY CAR	E CENTERS				
	Referral to te	ertiary care if		INTERVE		Descus	CONTEXT IN WHICH USEFUI		
				Psychoeducation			Poor adherence; high family expressed emotions		
				Family therapy Cognitive remediation			High family expressed emotions; family discord Poor neuro and social cognitive functions		
Inpati		n for clarification		Cognitive behavior therapy		Depression,	Depression, anxiety, obsessions, persistent psychotic symptoms		
	ory, thorough al status exami			Social skills training		Poor social	Poor social skills		
-	ostic psychomo or MRI, neurolo	etry, brain CT gy consultation		Vocational rehabilitation and supported education			Poor occupational functioning, challenges in studying or getting / pursuing gainful occupation		
and urine toxicology screen 2. Poor outcome: +Following psychosocial interventions may be offered in isolation or in combination depending on the context in inpatient, outpatient or day-boarding settings		1	Day care with interventions including vocational training, recreational activities, living-skill training, etc.		functioning	Negative symptoms, poor socio-occupational functioning, combination of other symptoms listed in the table			
		I	nterventions for	substance-use		use of substance or substan	nce use		
		Pi	Pregnancy – puerperium services Pregnancy and post-partum advise and interventions		interventions Pre-pregnand and post-partum advise an	су,			

ALGORITHM FOR CHOOSING ANTIPSYCHOTIC MEDICATION (AP) FOR TREATMENT OF SCHIZOPHRENIA

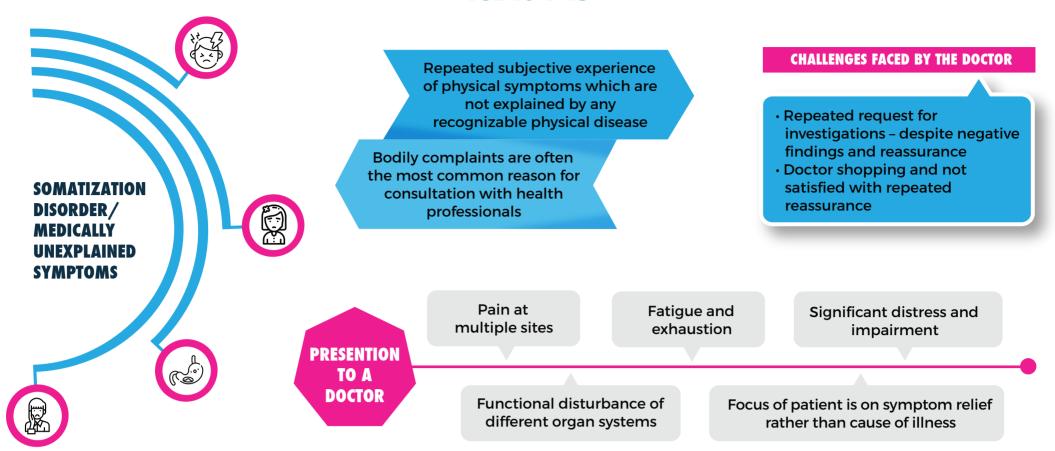


KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES





Standard Treatment Workflow (STW) for the Management of SOMATOFORM DISORDER (SD) ICD10-F45



DIAGNOSTIC CRITERIA

INITIAL ASSESSMENT

- Detailed clinical examination to rule out any medical illnesses which might explain the symptoms
- Complete history of the onset of all symptoms, exacerbating and relieving factors
- Assessment for any other psychiatric illness such as depression or anxiety disorders

PSYCHOSOCIAL ASSESSMENT

- · Encourage to talk about psychosocial stressors if any
- Individual factors poor copying skills, anxiety, life events, health anxiety, medical illnesses
- Family related factors Substance use in family, interpersonal relationship with family, financial status
- Environmental factors support system, peer relationship, work environment

DIAGNOSTIC CRITERIA

- A. One or more somatic symptoms that are distressing or result in significant disruption of daily life.
- B. Excessive thoughts, feelings, or behaviours related to the somatic symptoms or associated health concerns as manifested by at least one of the following:
 - 1. Disproportionate and persistent thoughts about the seriousness of one's symptoms
 - 2. Persistently high level of anxiety about health or symptoms
 - 3. Excessive time and energy devoted to these symptoms or health concerns
- C. Although only one somatic symptom may not be continuously present, the state of being symptomatic is persistent (typically more than 6 months)
 A persistent course is characterized by severe symptoms, marked impairment, and long duration (more than 6 months)

Following list include the commonest symptoms

- Pain symptoms at multiple sites (such as abdominal, back, chest, dysmenorrhea, dysuria, extremity, head, joint, rectal) is often present
- 2. Gastrointestinal sensations (pain, belching, regurgitation, vomiting, nausea)
- 3. Abnormal skin sensations (itching, burning, tingling, numbness, soreness) and blotchiness

Severity:

Mild - only one of the symptoms specified in criterion B is fulfilled

Moderate - Two or more of the symptoms specified in criterion B is fulfilled

Severe - Two or more of the symptoms specified in criterion B are fulfilled, plus there are multiple somatic symptoms (or one very severe somatic symptom)

4. Sexual and menstrual complaints (ejaculatory or erectile dysfunction, hyperemesis of pregnancy, irregular menses, menorrhagia, sexual indifference) are also common

MANAGEMENT

PRIMARY CARE

- Detailed physical examination
- Management of anemia and nutritional deficiencies
- Avoid irrational use of pain medications
- Low dose of antidepressant medications Amitriptyline 12.5 mg to 50 mg (max) night dose
- Explain that onset of medication effect will take 2-3 weeks
- Validate the somatic symptoms
- Advise to engage in routine activities, physical exercise and relaxation techniques like deep breathing
- Discuss with family members that the symptom, distress and disability are genuine
- Strengthen supports
- Regular follow up

TERTIARY CARE

- Inpatient care if needed
- Combination of two psychotropic medications (when required)
- Add on second and third line medications Duloxetine, Mirtazapine, anticonvulsants (Lamotrigine, Pregabalin). Use of Gabapentin, Carbamazepine if chronic pain symptom predominates
- Structured Cognitive Behavioural Therapy, Cognitive restructuring, Mindfulness and acceptance based approach
- Use of alternative medicine approach Yoga
- Collaborative approach involve Physician, Neurology team and Pain Clinic referral (where indicated)
- Vocational rehabilitation if needed
- Physical therapies guided exercise and physiotherapy



- 2. No improvement after 4 weeks of treatment with first line medications
- 3. Comorbid medical illness
- 4. Suicidal risk
- 5. Comorbid psychiatric illness

SECONDARY CARE

REFER TO

SECONDARY

CARE IF

- Investigations to rule out any medical illnesses that might explain the symptoms
- Complete history with behavioural observation
- Use 2nd line medications SSRIs (Escitalopram 10-20 mg, Sertraline 50-100 mg, Fluoxetine 20 mg) and SNRIs (Venlafaxine 75 – 150 mg, Duloxetine 30- 60 mg)
- Combination of two psychotropic medications (might be required if poor response to single medication)
- Brief counselling
- Psycho education focusing on relationship between stress and physical symptoms
- Relaxation training, regular exercise, yoga and meditation
 - 1. No improvement in 2nd line
 - treatment
 - 2. High suicidal risk

psychotherapy

4. Difficult patients

- 3. Needing intense counselling/
- REFER TO TERTIARY CARE IF

REFERENCES

Desai C & Chaturvedi SK. Medically Unexplained Somatic Symptoms & Chronic Pain – assessment & management. A primer for Healthcare professionals. 1st Edition 2017. Paras medical publisher, Hyderabad, India.

- World Health Organization. (2017). mhGAP training manuals for the mhGAP intervention guide for mental, neurological and substance use disorders in non-specialized health settings version 2.0 (for field testing). World Health Organization. http://www.who.int/iris/handle/10665/259161.
- Agarwal V, Srivastava C & Sitholey P. Clinical Practice Guidelines for the Management of Paediatric Somatoform disorders. Indian Psychiatric Society Practice guidelines 2018.
- Guidance for health professionals on medically unexplained symptoms (MUS) https://www.rcpsych.ac.uk/pdf/CHECKED%20MUS%20Guidance_A4_4pp_6.pdf
- · Jacob KS. A simple protocol to manage patients with unexplained somatic symptoms in medical practice. Natl. Med. J. India. 2004; 17: 326-8

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

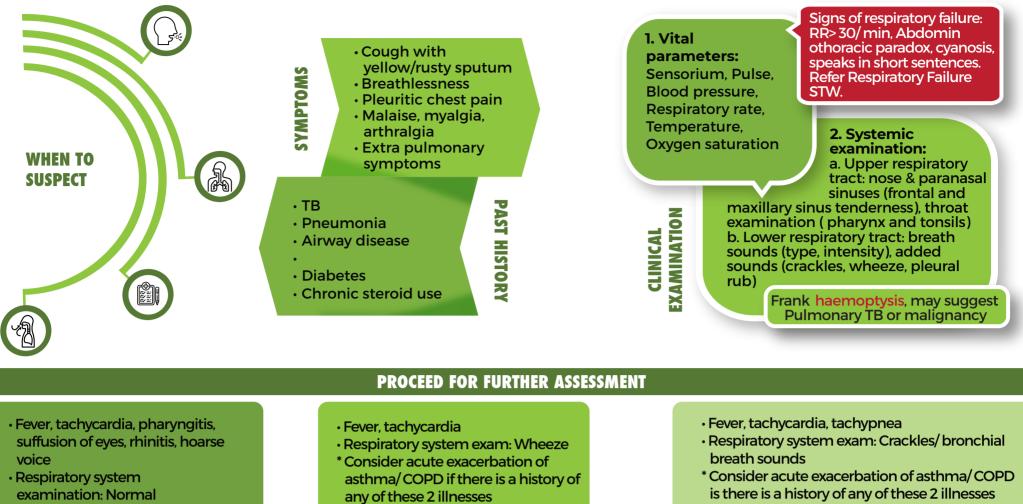


PULMONOLOGY





Standard Treatment Workflow (STW) for Management of ACUTE RESPIRATORY INFECTION IN ADULTS ICD-10-J09-J18; J00-06; J40



PATHWAYS BASED ON INITIAL ASSESSMENT FINDINGS

PATHWAY 1: ACUTE URI (RESPIRATORY CATARRH)

LABORATORY INVESTIGATION:

- Total and differential count in suspected flu. **TREATMENT**
- Symptomatic treatment for fever, myalgia (Paracetamol or other NSAID),
- Rest, Oral fluids (plenty)
- Oral antihistamines (Tab. CPM 4mg BD) for severe runny nose or sneezing
- Antibiotics in acute follicular tonsillitis: Amoxicillin/Ampicillin 500mg tid X 5 days In penicillin sensitive individuals: Erythromycin estolate 250mg q 6 hrly X 5
- days with food Suspect epidemic flu

H/ o recent travel, symptoms of upper respiratory infection, diarrhoea, myalgia, breathlessness **Refer to higher centre for diagnosis, notification and treatment.**

PATHWAY 2: ACUTE BRONCHITIS

LABORATORY INVESTIGATION:

- Total and differential count if sputum is purulent,
- X-ray chest PA view
- TREATMENT
- Symptomatic treatment for fever (Paracetamol or other NSAID), Oral fluids (plenty)
- Inhaled bronchodilators: Salbutamol nebulization (5mg/2.5ml) 6-8 hourly
- Antibiotics if there is purulent sputum and
- polymorphonuclear leukocytosis
 Amoxicillin 500mg tidX 5 days
 - In penicillin sensitive individuals: Erythromycin estolate 250mg q 6 hrly X5 days with food
- If asthma is suspected refer to asthma STW

PATHWAY 3: COMMUNITY ACQUIRED PNEUMONIA

SEVERITY ASSESSMENT

- X-ray
- Use CRB-65* score for mortality risk assessment in primary care

CRB-65 SCORE						
SCORE	RISK CLASS	SITE OF CARE				
0	Low Risk	OP				
1-2	Intermediate Risk	IP				
3-4	High Risk	ICU				

*65 in the scoring mnemonic refers to age>65

- Give 1 point for each of the following Prognostic features: • Confusion
- Respiratory rate ≥30/ min
- Low BP (DBP ≤ 60 mm Hg or SBP ≤ 90 mm Hg)

TREATMENT

Age ≥65 years

INVESTIGATIONS

Preliminary

- Chest radiogram
- Repeat if:
- i. Patient is not improving/worsening clinically
- ii. Suspected underlying malignancy **Desirable**
- 1. Pulse oximetry in outpatients
- 2. Sputum microbiology: In suspected PTB & non-response after 48 hours of antibiotics
- 1. Targeted towards Streptococcus pneumoniae
- 2. Oral antibiotics after checking for comorbidities* (Diabetes, CVDs, CKD, CLD, Hepatic Pathology, Cancer, Alcohol Abuse, H/ o antibiotics within last 3 months.)
 - a. Without comorbidities: Cap. Amoxicillin (500 mg TDS)/Tab. Erythromycin 250mg QID/Tab. Doxycycline 100mg BD
 - b. With comorbidities: Cap. Amoxicillin 500mg TDS + Tab.Azithromycin 500 mg OD
- 3. Duration: 5 days in (A); extend to a 7-10 days course if there is no response within 3 days of starting treatment and in (B).
- 4. Do not give:

a. Corticosteroids: unless other medical indications present b. Fluoroquinolones: as they have anti-tubercular activity.

INPATIENT MANAGEMENT OF CAP

ANTIBIOTIC THERAPY IN THE HOSPITALIZED NON-ICU SETTING

a. Single agent IV β -lactam

b. If suspected atypical pathogens, other end organ disease, diabetes, malignancy, severe CAP, use of antibiotics in past 3 months: Combination of IV β -lactam (Cefotaxime 2 grams TID/IV Ceftriaxone 1gram BD/Amoxicillin–Clavulanic acid 1.2 grams TID) + ORAL macrolide (Tab Azithromycin 500 mg PO OD/Tab Clarithromycin 500 mg PO BD)

ANTIBIOTIC THERAPY IN THE HOSPITALIZED ICU SETTING

i. Patients without risk factors for Pseudomonas aeruginosa: Manage as above ii. Suspected P. aeruginosa (diabetes, chronic lung disease like bronchiectasis, chronic steroid therapy):

IV Cefepime (IG BD)/ IV Ceftazidime (2G TID)/ Piperacillin-tazobactam(4.5 G QID)/ IV Cefoperazone-sulbactam 1.5G IV TID/ IV Meropenem 1g TID; Combination therapy : Aminoglycosides(IV Amikacin)/ Antipseudomonal

fluoroquinolones(Levofloxacin/Moxifloxacin)

REFERRAL TO A HIGHER CENTRE : CLINICAL CRITERIA

- 1. Frank hemoptysis and /or Signs of respiratory failure [listed under in the history and evaluation sections]
- 2. CRB-65 score > 1
- 3. Oxygen saturation by pulse oximetry \leq 92% (patients \leq 50 yrs) OR <90% (patients > 50 yrs)
- 4. Multi-lobar consolidation on chest X-ray
- 5. Confusion/disorientation
- 6. Hypothermia (core temperature<360C)

ADJUNCTIVE THERAPIES FOR THE MANAGEMENT OF CAP

a. Steroids are not recommended for use in non-severe CAP b. Non-invasive ventilation may be used in patients with CAP and acute respiratory failure

CONTRA INDICATIONS FOR NON-INVASIVE VENTILATION a. Cardiorespiratory arrest

b. Presence of severe upper airway inflammation & edema

- c. Severe haemodynamic instability hypotension
- d. Eu-capnic (normal PaCO2) coma
- e. Multiple organ dysfunction or severe psychomotor agitation

DISCHARGE CRITERIA

Accepting orally, Afebrile and Hemodynamically stable for a period of at least 48 h

POINTS TO NOTE WHILE SHIFTING

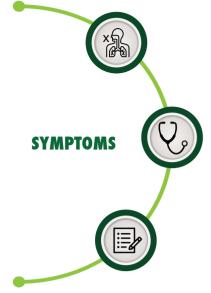
- If referring to a higher center, give the first dose of antibiotic (oral and if available, parenteral), secure an IV line and start 0.9% Normal saline and oxygen supplementation through face mask at 4-6 litres per minute during shift
 If the patient is drowsy, has copious secretions, consider
- calling for help from the SUB-DISTRICT/DISTRICT hospital for endotracheal intubation and shifting on a transport ventilator

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES





Standard Treatment Workflow (STW) for the Management of ASTHMA ICD-10-J45



- Classic symptoms
 Recurrent/episodic wheezing
- Breathlessness
- Cough and/or chest tightness

Supportive features

- History of atopy, family history of asthma, presence of triggers, presence of rhonchi on chest auscultation
- No alternative explanation for
- these symptoms

TRY AND RULE OUT

- Other obstructive airway disorders - see Table 1 for features that favour asthma over COPD
- Other mimics presence of fever, constitutional symptoms, purulent sputum, hemoptysis, focal chest signs on physical examination, foreign body aspiration, abnormal chest radiograph, etc.

APPROACH TO DIAGNOSIS

- Clinical assessment is the mainstay
- Airway obstruction, and bronchodilator reversibility, on
- spirometry (if available) may support diagnosis
- Refer patients for further work-up if diagnosis is in doubt

INITIATION AND MODULATION OF ASTHMA PHARMACOTHERAPY

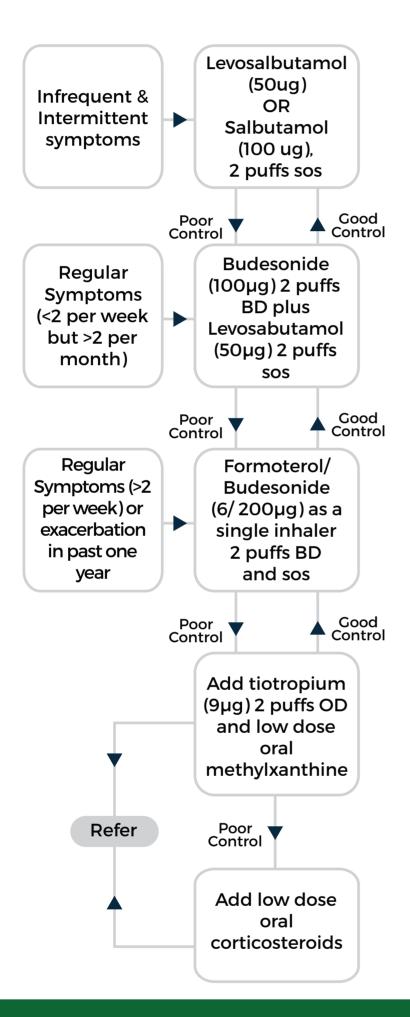


TABLE 1. DIFFERENTIATING BETWEEN ASTHMA AND CHRONIC OBSTRUCTIVEAIRWAY DISEASE (COPD)

	Asthma	COPD
Age of Onset	More often in childhood or early adulthood; variable	Usually later in life (4th or 5th decade)
Course	Episodic	Progressive
Smoking, other exposures	Uncommon	Common
Nasal Symptoms, Atopy	Common	Rare
Family History	Often	Uncommon
Triggers	Often Identified	None
Wheeze	Prominent and almost universal	May or may not be present

TABLE 2. LEVEL OF CURRENT ASTHMA CONTROL (OVER THE PRECEDING FOUR WEEKS)

Components	Inadequately controlled (any one)	Adequately controlled (all should be present)
Daytime symptoms or use of rescue medication	More than twice a week	Twice or less in a week
Night-time symptoms/ awakening	Any	None

Limitation of activities	Any	None
Pulmonary function (if available)	FEV1 <80% of predicted or PEF <80% of personal best	FEV1 >80% of predicted or PEF >80% of personal best

FEV1 Forced Expiratory Volume in first second, PEF Peak Expiratory Flow

GUIDING PRINCIPLES

- Mainstay of pharmacotherapy: Inhaled drugs
- Frequency of symptoms determine treatment initiation (see figure 1 for details)
- Reassess at 3-4 weeks good response: in favour of asthma diagnosis
- Patient education for compliance, warning signs, triggers, inhaler technique, PEF monitoring
- Inhaler technique to be monitored
- Follow-up at 4-12 weeks, assess diseases control by clinical parameters (see Table 2)
- · Step-up or step-down treatment as per level of asthma control (see figure 1)
- · Follow up three-monthly and modulate treatment as needed
- Refer for further evaluation and management if asthma remains poorly controlled

DISEASE EXACERBATION

WHEN TO SUSPECT EXACERBATION

- Suspect if acute symptomatic worsening, or reduction in PEF to below 80% of personal best, while on continued treatment
- Take two additional puffs of the inhaler used if symptoms persist, and repeat if needed
- If no response after 24 hours, or symptomatic worsening, or further reduction in PEF, contact physician
- Physician to assess severity of exacerbation and manage accordingly

LIFE-THREATENING EXACERBATION

Altered sensorium, orthopnea, cyanosis, paradoxical breathing, hypotension, and/ or bradycardia (heart rate <60 bpm) – immediately refer to higher centre with ICU facility

SEVERE ACUTE ASTHMA (PATIENT TO BE ADMITTED)

- Inability to complete sentences, agitation, use of accessory muscles, respiratory rate >30/ min, heart rate >110/ min, pulsus paradoxus >25 mm Hg, silent chest, and/ or room air sPo2 <92%
- Oxygen supplementation to maintain spO2 92-95%
- Nebulized levosalbutamol/ipratropium (1.25 mg/ 0.5 mg) three doses at 20-minute interval, then 4-6 hourly or as needed
- Injection hydrocortisone 200 mg intravenously, then oral prednisolone 0.5 mg/ kg daily for five days
- Refer if no improvement
- Dischargeonly when symptoms improve, wheezing absent or significantly reduced, heart rate <100 bpm, respiratory rate <30/ min, room air sPo2 >94%
- Schedule follow-up outpatient visit at one week

NON-SEVERE ACUTE ASTHMA

- If none of the above features present manage on outpatient basis
 - Continue additional inhaler doses as needed
 - Oral prednisolone 0.5 mg/ kg daily for five days
 - Schedule follow-up outpatient visit at one week

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

REFERENCES

- 1. Agarwal R, et al. Guidelines for diagnosis and management of bronchial asthma: Joint ICS/ NCCP(I) recommendations. Lung India 2015;32(Suppl 1):S3-S42.
- 2. Clobal Initiative for Asthma (GINA). Global strategy for asthma management and prevention. 2018.
- 3. National Institute for Health and Care Excellence (NICE). Asthma: diagnosis, monitoring and chronic asthma management. 2017.

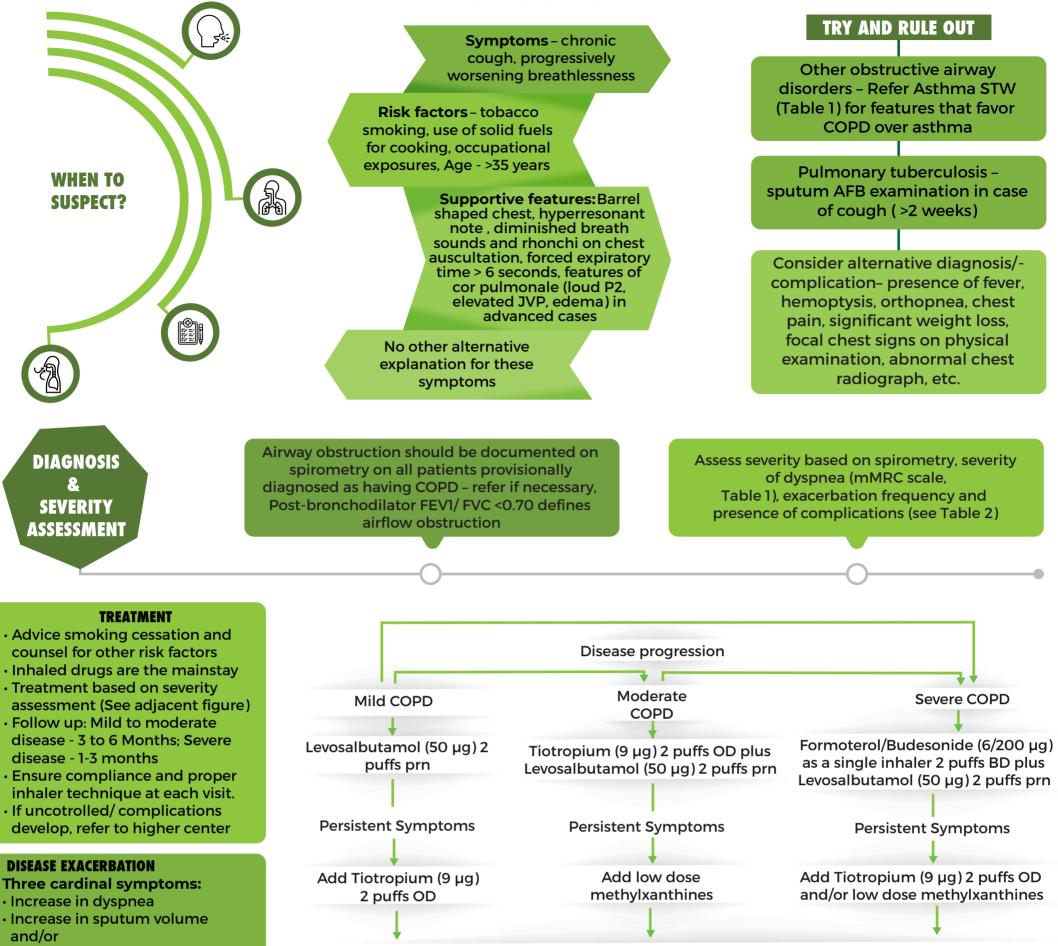






Standard Treatment Workflow (STW) for the Management of CHRONIC OBSTRUCTIVE PULMONARY DISEASE

ICD-10-J44.9



Increse in sputum purulence

Classify As:

- Mild Exacerbation
- Severe Exacerbation

Features Of Severe Exacerbation:

- Cyanosis
- Respiratory rate >30/ min
- Heart rate >110/min
- Systolic blood pressure <90 mm Hg
- SpO2 <90%
- Paradoxical respiratory movements
- Altered sensorium
- Asterixis
- Presence of severe co-morbid conditions (e.g. heart failure, arrhythmia)

MILD EXACERBATION

- Increase dose and/ or frequency of levosalbutamol and/ or ipratropium inhalation, or nebulized levosalbutamol/ ipratropium (1.25 mg/ 0.5 mg), repeated as needed at 20-minute interval
- Amoxycillin 500 mg TDS/ Azithromycin 500 mg OD/ Doxycycline 100 mg OD (BD on day 1) X 5 Days
- Oral prednisolone 30 mg daily X
 5 days

SEVERE EXACERBATION

Treatment as under Mild Exacerbation

Supplement oxygen with target spO2 of 92% (if spO2 monetoring available)

Refer if inadequate response, onset of new complications, or suspicion of alternative diagnosis

TABLE 1. GRADING OF BREATHLESSNESS USING MODIFIED MEDICAL RESEARCH COUNCIL (MMRC) SCALE.

GRADE	DESCRIPTION OF BREATHLESSNESS
0	I only get breathless with strenuous exercise.
1	I get short of breath when hurrying on level ground or walking up a slight hill.
2	On level ground, I walk slower than people of the same age because of breathlessness or have to stop for breath when walking at my own pace.
3	I stop for breath after walking about 100 yards or after a few minutes on level ground.
4	I am too breathless to leave the house or I am breathless when dressing.

TABLE 2. SEVERITY CLASSIFICATION FOR COPD

SEVERITY	POSTBRONCHODILATOR FEV1 (% PREDICTED)	DYSPNEA (MMRC GRADE)	EXACERBATIONS IN LAST ONE YEAR	COMPLICATIONS*
MILD	<u>≥</u> 80	<2	<2	NO
MODERATE	50-79	<u>≥</u> 2	<2	NO
SEVERE	<50	<u>≥</u> 2	<u>≥</u> 2	YES

The category with the worst value should be used for severity classification *Complications include respiratory failure, cor pulmonale, and secondary polycythemia

RED FLAG SIGNS FOR PEOPLE HAVING EXCERBATION

- Altered sensorium
- spO2 <88% despite therapy
- Heart rate >110 bpm
- Systolic blood pressure <90 mm Hg
- High risk comorbid conditions (arrhythmia, congestive cardiac failure, poorly controlled diabetes, renal or liver failure)

Refer to higher centre for further management, and ensure continued supplemental oxygen and nebulization during transfer

SCHEDULE FOLLOW UP VISIT ONE WEEK AFTER DISCHARGE

ADMISSION CRITERIA

- 1. Severe symptoms; sudden worsening of resting dyspnea,
- 2. Fall in oxygen saturation, cyanosis, confusion, drowsiness.
- 3. Failure of an exacerbation to respond to initial medical management.
- Presence of serious comorbidities (heart failure, newly occurring arrhythmias, etc.)

DISCHARGE CRITERIA

- 1. Normalization of clinical and laboratory data to pre-admission levels
- 2. Patient able to follow maintenance therapy
- 3. Completion of acute medications
- 4. Adequate control of comorbidities

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES

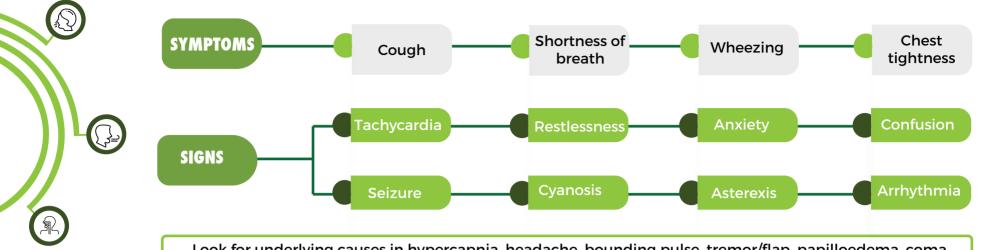
REFERENCES

- 1. Gupta D, et al. Guidelines for diagnosis and management of chronic obstructive pulmonary disease: Joint ICS/ NCCP(I) recommendations. Lung India 2013;30:228-67
- 2. Global Initiative for Chronic Obstructive Lung Disease (GOLD). Global strategy for the diagnosis, management, and prevention of chronic obstructive pulmonary disease. 2019 report.
- 3. National Institute for Health and Care Excellence (NICE). Chronic obstructive pulmonary disease in over 16s: diagnosis and management. 2018.





Standard Treatment Workflow (STW) for the Management of **RESPIRATORY FAILURE** ICD 10: J96.0



Look for underlying causes in hypercapnia, headache, bounding pulse, tremor/flap, papilloedema, coma.

ARI

LRTI

embolism

HYPOXIA (SPO2 <90%) PNEUMONIA/ LRTI **PULMONARY EMBOLISM HEART FAILURE SYMPTOMS** SIGNS **SYMPTOMS** SIGNS **SYMPTOMS** SIGNS Dyspnea or Tachycardia • Cough with or Tachypnea Sudden Shortness • Syncope exertion or rest Pulsus Alterans without Sputum Tachycardia • Arrhythmia of Breath Crackles and Rhonchi • Weak Rapid Thready Pulse Chest Pain Chest Pain Chest Pain Tachycardia Wheezing Pink Frothy Sputum • Fever with Chills, Hypoxemia • Calf Pain & or Pleuritic Chest Pain Fatigue Cyanosis Fatigue, Malaise Swelling Pallor Hemoptysis Distended Neck Veins **AIRWAY DISEASE ACUTE ASTHMA AE OF COPD BRONCHIOLITIS SYMPTOMS SYMPTOMS** SIGNS SIGNS **SYMPTOMS** SIGNS Wheeze • Tachypnea • Worsening of Tachypnea • Cough Cyanosis Tachycardia Shortness of Nasal Flares Shortness of Hypoxemia Dyspnea **Breath** • Fall in SPO2 Increase in Hypercarbia **Breath** • Tachypnea Paradoxical Breathing Chest Tightness Use of Accessory Sputum Confusion Wheezing • Cough Muscle Production Drowsy (children) Increased Cough Peripheral Edema Crackles and or Rattling sounds in Lung **INVESTIGATIONS** Spirometry(COPD, Neuromuscular disease **Chest Xray** Sputum culture, Blood culture (if febrile) ABG, CRP, FBC, U&E TREATMENT Acute Severe Heart Pneumonia Pulmonary

AE COPD

DIAGNOSIS

failure

Asthma

OXYGEN	Start oxygen therapy at SpO2 < 90% Monitor SpO2 during oxygen therapy to titrate flow rate: target SpO2 < 96% Oxygen delivery usign Nasal cannulae/ Simple face mask/ Venturi mask/ Non re-breathing mask (Note: for patients with AECOPD, keep lower target SpO2 = 88-92%)							
BRONCHODILATORS	SOS	SABA ± SAMA (Salbutamol ± Ipratropium neb q20 min X 1 hr then prn)	SABA + SAMA (Salbutamol neb hourly + Ipratropi- um neb 4 hourly)	SABA + SAMA	SOS	SOS		
DIURETICS	Yes (IV Furosemide 40 mg or Torsemide 20 mg)	SOS	SOS	SOS	SOS	SOS		
ANTIBIOTICS			No risk factor Pseudomonas: Ceftriaxone or levofloxacin or moxifloxacin Pseudomonas risk factor: levofloxacin or piperacillin tazobactam or ceftazidime or cefepime Influenza suspect: Oseltamivir		Mild/ Mod cases: Amoxycillin PO/ IV or Ceftriaxone IV Severe Cases: Amoxycillin IV or Ceftriaxone IV Atypical pneumonia: Azithromycin IV/ PO or Doxycycline IV/ PO			
STEROIDS		Yes (Methylpredniolone IV 40 to 60 mg or Prednisolone PO 60 mg)	Yes (Methylprednisolone IV 60 to 125 mg IV q6-12 hourly)	Yes	Severe CAP (fiO2 > 0.5 AND pH <7.3 OR lactate >4 mmolL-1 OR CRP > 150 mgL-1): Methylprednisolone IV 0.5 mg/ kg q12h			
LMWH	Prophylactic, if indicated	Prophylactic, if indicated	Prophylactic, if indicated	Prophylactic, if indicated	Prophylactic, if indicated	If high suspicion with low risk of bleeding: UFH (if thrombolysis anticipated), OR LMWH		
REFERRAL	REFERRAL No relief OR Need for mechanical ventilation OR life threatening features: Stabilize CAB, transfer to higher center							
			ABBREVIATIONS					
• LRTI : Lower Respiratory Tract Infection • LMWH:Low Molecular Weight Heparin• SABA : Short Acting Beta Agonist • SAMA: Short Acting Muscarinic Antagonist• CAP: Community Acquired Pneumonia • UFH : Unfractionated Heparin								
		🖝 KEEP A HIGH	THRESHOLD FOR INVASIVE	PROCEDURES				
This STW has been prepared by national experts of India with feasibility considerations for various levels of healthcare system in the country. These broad guidelines are advisory, and								

are based on expert opinions and available scientific evidence. There may be variations in the management of an individual patient based on his/her specific condition, as decided by the treating physician. There will be no indemnity for direct or indirect consequences. Kindly visit our web portal (stw.icmr.org.in) for more information. © Indian Council of Medical Research and Department of Health Research, Ministry of Health & Family Welfare, Government of India.



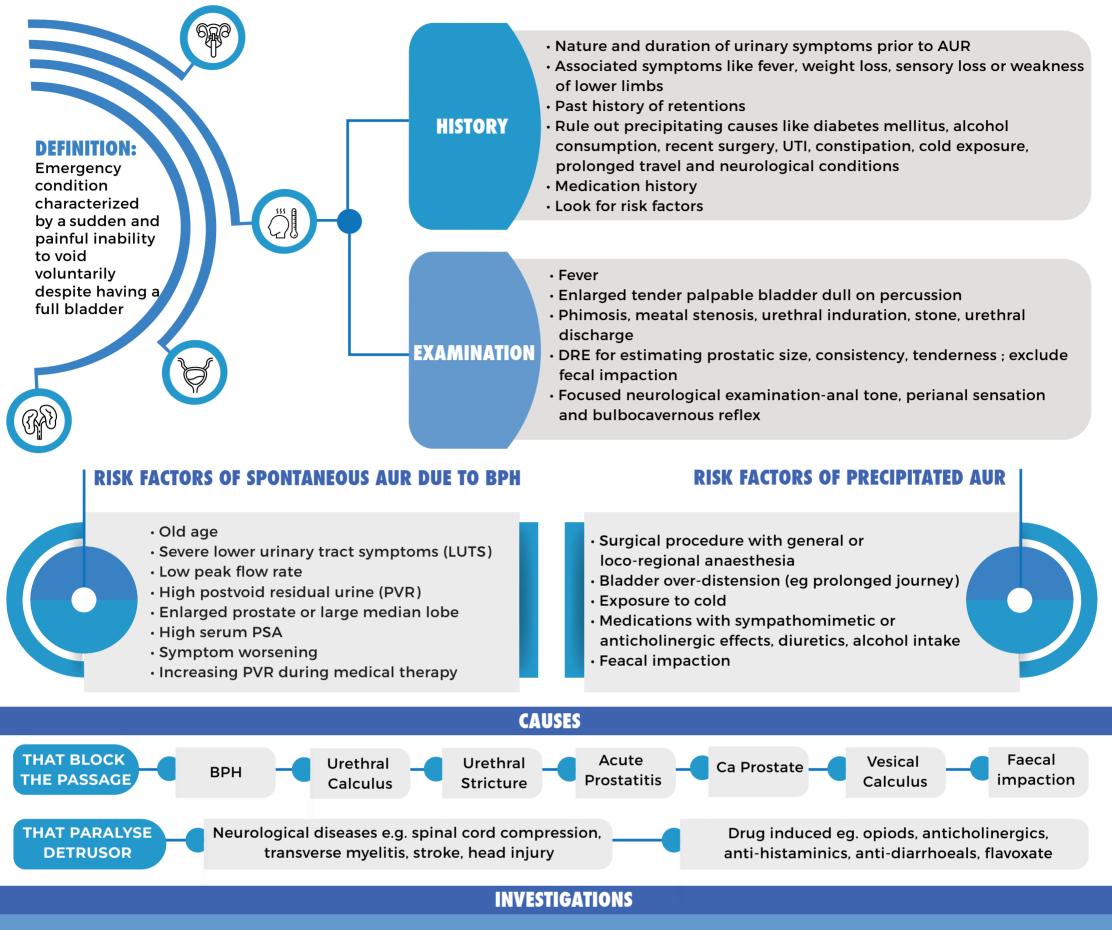






Standard Treatment Workflow (STW) for the Management of ACUTE URINARY RETENTION IN MEN (AUR)

ICD-10-R33.9



As AUR is an acute emergency, no investigation is required before catheterization to relieve symptoms. The volume of urine

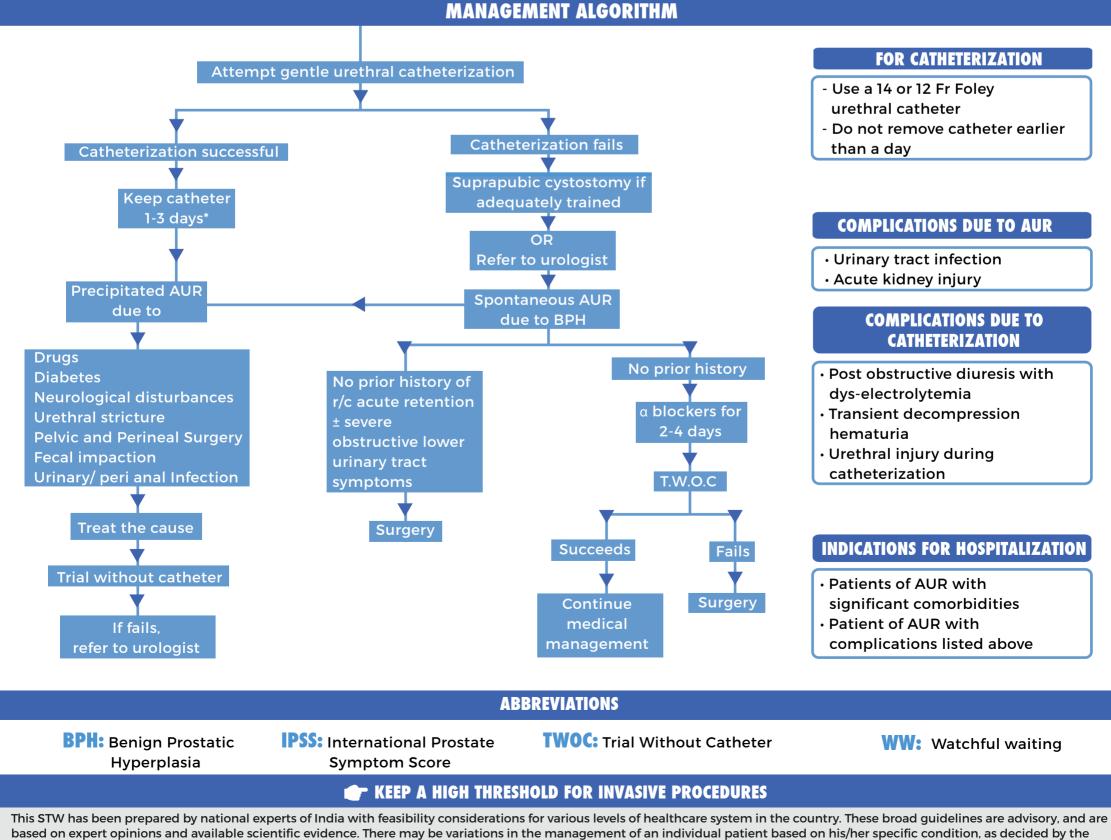
drained should be documented.

DESIRABLE

CBC, S. Glucose, S. Creatinine and Electrolytes, USG KUB Urine analysis& Urine culture of the drained urine

OPTIONAL (ONLY BY SPECIALISTS) NOT TO BE DONE ROUTINELY

• Cystoscopy,CT / MRI,RGU + MCU,Urodynamic studies



treating physician. There will be no indemnity for direct or indirect consequences. Kindly visit our web portal (stw.icmr.org.in) for more information.

Indian Council of Medical Research and Department of Health Research, Ministry of Health & Family Welfare, Government of India.

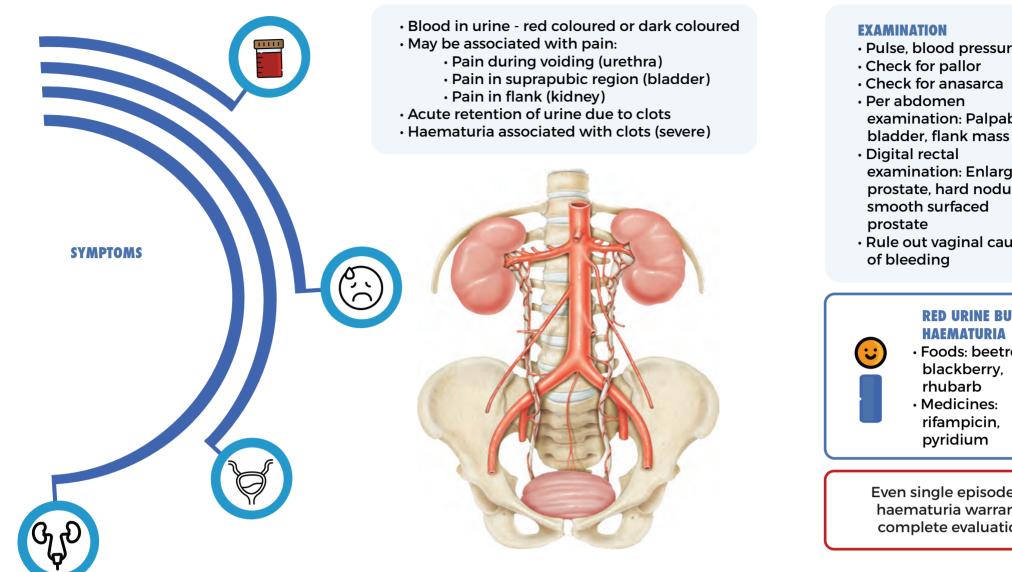




Standard Treatment Workflow (STW) for the Management of **GROSS HAEMATURIA**

ICD-10-R31.0

PERFORM THOROUGH CLINICAL EVALUATION



MAKE A CLINICAL DIAGNOSIS: IS HAEMATURIA

INITIAL

• Urethra: stone, urethritis, stricture Prostate: inflammation, benign hyperplasia, malignancy

TOTAL

· Kidney: stone, malignancy (renal parenchyma, pelvis/ ureter), genito-urinary tuberculosis

HOW TO INVESTIGATE

ESSENTIAL

- Urine examination routine, microscopy
- Hemoglobin estimation
- Kidney function tests (KFT)
- Ultrasonography of kidney urinary bladder and prostate region

DESIRABLE

 Contrast enhanced computed tomography of kidney urinary bladder region/ intravenous pyelography (if KFT normal)

OPTIONAL

- Urine culture • Urine for active sediments(if nephrotic/ nephritic syndrome suspected) PT/INR (if
- bleeding disorder suspected)
- Serum prostate specific antigen (if required)

- Pulse, blood pressure
- examination: Palpable
- examination: Enlarged prostate, hard nodular/
- Rule out vaginal causes

RED URINE BUT NOT

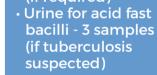
Foods: beetroot,

Even single episode of haematuria warrants complete evaluation

- Ureter: stone, malignancy, genito- urinary tuberculosis
- Bladder: infection, genitourinary tuberculosis, stone, malignancy

TERMINAL

- Bladder: stone, tumor at bladder neck Prostate: inflammation, benign
- hyperplasia, malignancy
- Magnetic resonance imaging of Kidney urinary bladder region (if KFT deranged)
- Urine cytology if > 40yrs or smoker Cystoscopy if > 40 years or smoker



WHEN TO REFER (WARNING SIGNS) Deranged kidney functions

- Suspecting malignancy
- Haematuria with hypertension / albuminuria
- · Persistent severe haematuria

HOW TO TREAT

GENERAL

- Start intravenous fluids if required (primary level)
- · If Anaemia may transfuse blood as required (primary level)
- Manage clot colic / flank pain with analgesics (primary level)
- · If Acute urinary retention - catheterise with 20/22Fr 3 way Foley and may start continuous irrigation with normal saline (Primary level)
- Cystoscopic clot evacuation may be performed if feasible (tertiary level)
- If basic evaluation and management facilities are unavailable - refer (tertiary level)

· Haematuria should be considered as a symptom of genitourinary malignancy in patients >40years old until proven otherwise

SPECIFIC

- Suspected nephrotic/nephritic syndrome: cola coloured urine, proteinuria, anasarca, hypertension Refer to nephrologist (tertiary level)
- Suspect urinary tract infection : presents with dysuria, increased frequency of voiding and other irritative lower urinary tract symptoms with/ without fever- treat with broad spectrum oral antibiotics (primary level)

DIFFERENTIAL DIAGNOSIS FOR CHRONIC CONDITIONS LEADING TO HAEMATURIA						
	Stones	Renal cell cancer	Bladder tumor	Genito-urinary tuberculosis		
Symptoms	Flank pain Ureteric colic Recurrent urinary tract infection Haemturia	Flank mass Flank pain Haematuria	Haematuria Urinary retention	Dysuria Frequency Nocturia Haematuria		
Investiga- tions	Ultrasonography Xray KUB Intravenous pyelography or Computed tomography	Ultrasonography Computed tomography	Ultrasonography Computed tomography Urine cytology	Urine analysis Urine acid fast bacilli Urine tuberculosis culture Gene expert (optional) Intravenous pyelography or Computed tomography		
Treatment	>5mm or symptomatic - refer to urologist	Mostly surgical treatment - refer to urologist	Mostly surgical treatment - refer to urologist	Oral Antitubercular treat- ment - 6months, refer to a urologist, close follow up		

REFERENCES

1. Standard treatment guidelines in urology: Ministry of Health and Family selfare

KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES





Standard Treatment Workflow (STW) for the Management of MALE INFERTILITY

ICD-10-N46.9

	HOW TO PROCEED?	AIM
WHEN TO SUSPECT?	Both partners examined simultaneously*	 To ascertain contributory male factor Identify potentially correctable conditions Identify incorrectable conditions
WHEN TO SUSPECT? Inability to conceive even after one year of regular unprotected intercourse. Evaluation earlier than one year if female age is >35yrs, family history of infertility or very anxious couples. Infertility Incidence is 10-15%. Male factor-contributory in 50% cases.	<text></text>	 Identify incorrectable conditions that may or may not be amenable to Assisted Reproductive Technique (ART) Identify underlying medical conditions responsible for infertility PHYSICAL EXAMINATION Body habitus (obesity, Klinefelter's). Secondary sexual characters, gynecomastia Penis: hypospadias, epispadiasis, chordee, Testes: volume, consistency, masses, contours Epididymis: flat, turgid, nodularity. Vas deferens -present/absent thickened or beaded Cords-presence of varicocele. Inguinal or scrotal scar. Rectal examination: cyst, dilated
	the chance of success.	seminal vesicles.

HISTORY

- Age of partners and duration of infertility.
- Use of contraception and lubricants.
- Knowledge of sexual cycle, technique frequency
- technique, frequency. • Sexual and ejaculatory dysfunction, volume of ejaculate
- Medical illness: STD, diabetes, recent fever, chronic bronchitis and any debilitating medical conditions
 - H/o Chemotherapy, Radiotherapy
 Congenital anomalies,
 - cryptorchidism, hypospadias, Chordee
 - Testicular torsion, drug history, trauma and swelling
 - H/o past surgeries(hernia repair, orchiopexy, retroperitoneal surgery)
 - Family history (infertility,consanguinity,genetic disorders),
 - Exposure to environmental toxins (pesticides,herbicides, chronic heat and radiation (sauna bath, tight non cotton undergarments, laptops & mobile)
 - Partner history: Any menstrual abnormality, infertility evaluation till date

INVESTIGATIONS

SEMEN ANALYSIS (ESSENTIAL)

- At least two- samples 1-2 months apart ; Abstinence of 1-3 days.; Collected in sterile, medical grade plastic wide mouth containers.
- Provided within the lab or transported within an hour at room temperature and examined immediately
- WHO 2010 criteria for normal report. Volume: >1.5, ml, Sperm conc.: >15 million/ml, Sperm motility: >40%, Progressive > 32%, Sperm morphology: >4% normal forms, Leukocyte density: <1 million/mL

DIAGNOSTIC CATEGORIES ACCORDING TO SEMEN ANALYSIS REPORT

- Normal Semen Analysis: Rule out sexual dysfunctions, Anatomic abnormalities, Female factor and unexplained
- Low volume semen: Incomplete Collection, Retrograde ejaculation, Ejac. duct obstruction, Cong. Absence of VasDeferens, Hypogonadism
- Azoospermia:
- Obstructive (Epididvmal.vasal)
- (Epididymai,vasai)
 Nonobstructive: (Genetic, Chromosomal, Hormonal, CT/RT, Post torsion testes, orchitis, Cryptorchidism, Idiopathic)
- Oligo-astheno-teratospermia: Isolated Asthenospermia: Antisperm antibodies, Sperm structural defect, Hypogonadism
- Multiple defects: Varicocele, Cryptorchidism, Genital tract infection, Systemic illness, Prolonged abstinence, Drugs (Sulfasalazine, NFT, Colchicine, Chemotherapy, GnRh analogs, Spironolactone, Ketokonazole, Anabolic steroids, cocaine, alcohol. Chemicals: heavy metals, herbicides, organic solvents, fungicides, pesticides)

Note: If a patient is unable to produce semen consider retrograde ejaculation and anejaculation. Need further evaluation.

OPTIONAL INVESTIGATIONS

- Hormonal assay: Serum FSH, LH, Prolactin, Testosterone, Estradiol, T/E ratio
- Culture: Urine, Semen, Prostatic fluid, Antisperm antibodies, Viability assay, Sperm function tests, Scrotal USG & doppler, TRUS, Genetic studies,
- Testicular biopsy (Multiple bilateral preferable)

MANAGEMENT

TREATMENT ALGORITHM

PHC/CHC

History and Physical examination(PE)

DISTRICT HOSPITAL Hormonal assay and Testicular biopsy

Proper Semen analysis

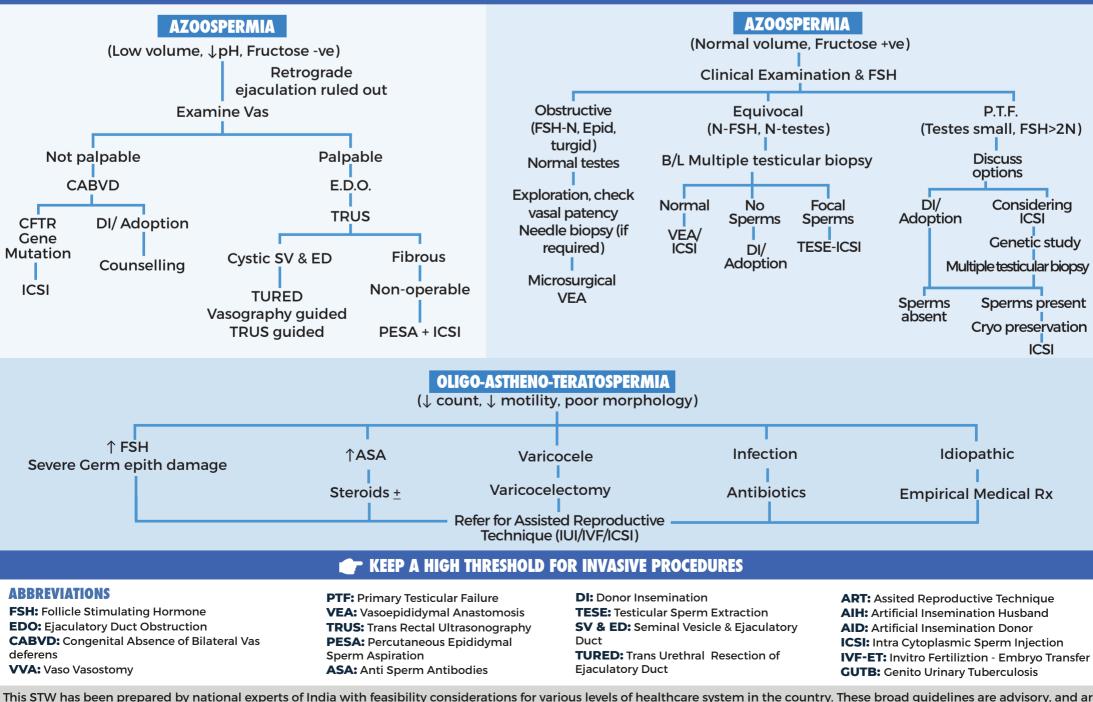
Normal Semen report: (Rule out unconsummation, sexual dysfunction, anatomic abnormailities) Abnormal Semen report:

- Refer to Urologist/infertility centre
- Preventive measures: Avoid gonadotoxins, gonadotoxic drugs, smoking, tobacco, chronic heat, excess use of mobiles; Encouraging healthy life style: Nutritious diet, regular physical exercise, avoid stress, use of antioxidants and vitamins(Vit. C, Vit E, Zinc)
- \cdot Female partner to be evaluated by gynecologist
- Management of reversible nonsurgical causes (Infections etc.) and surgical cause i.e. varicocoele if surgeon available.
- · For further evaluation refer to district/tertiary hospital.

- Management of sexual and ejaculatory dysfunction
- Management of Varicocele and Hypogonadotropic hypogonadism
- ART: AIH/AID and counselling for adoption.

TERTIARY LEVEL

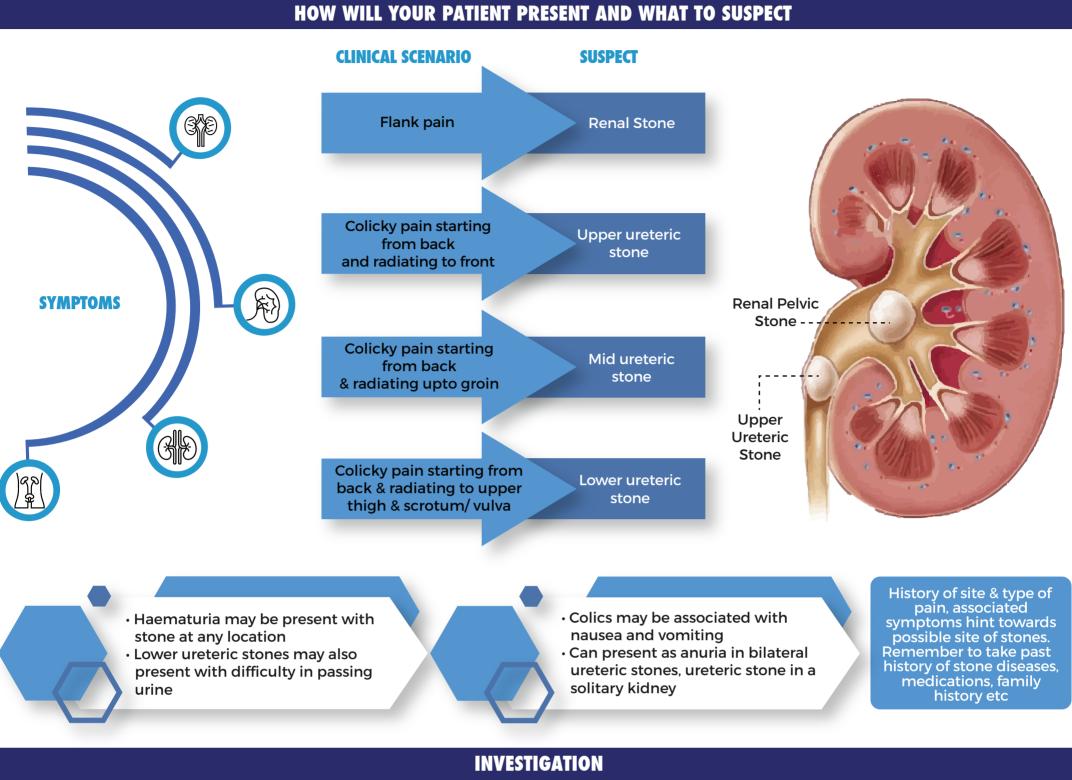
- Additional testing:TRUS, Genetic, ASA, Sperm function tests
- Advanced surgery: Microsurgical VVA,VEA, Varicocelectomy, TURED, Sperm retreival techniques, Cryopreservation and sperm banking
 Advanced ART: IVF-ET/IVF ICSI







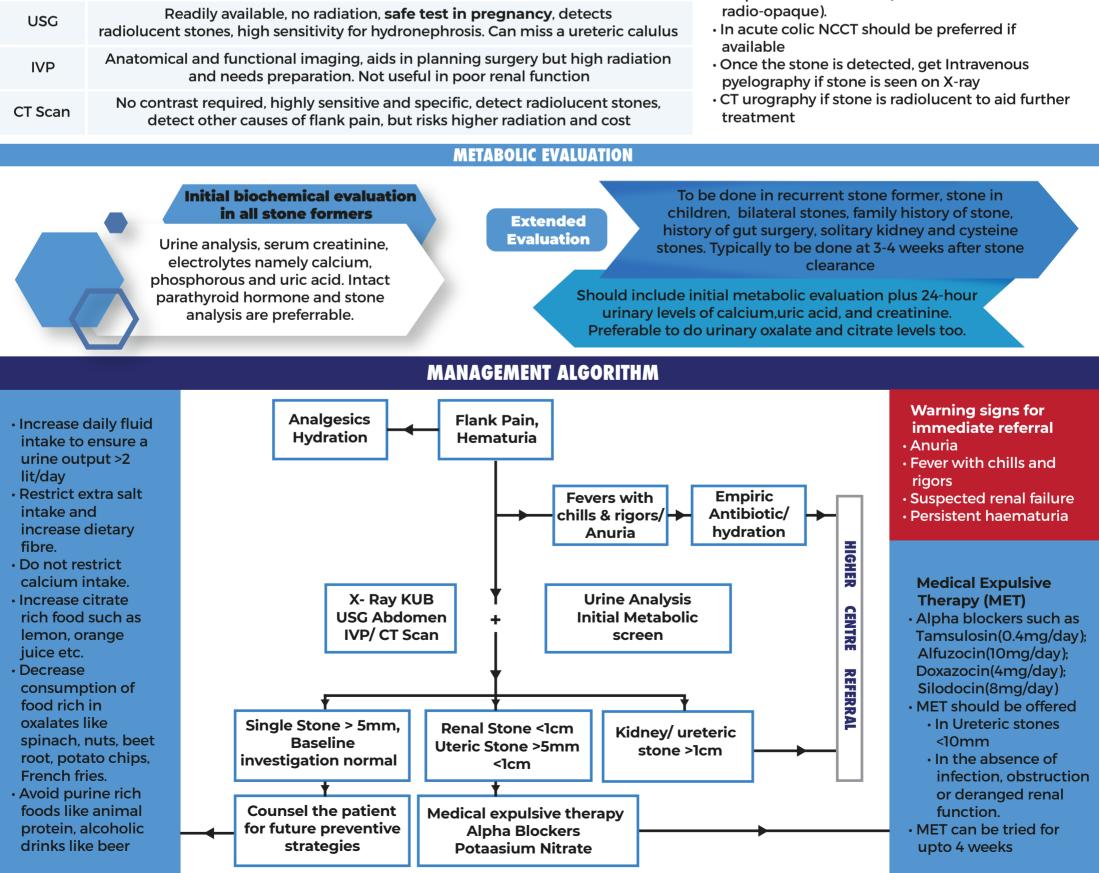
Standard Treatment Workflow (STW) for the Management of **RENAL AND URETERIC STONES** ICD N20.0



NAME	ADVANTAGES AND DISADVANTAGES
X-KUB	Readily available, inexpensive, minimal radiation but needs preparation hence may not be the preferred test in emergency settings

TIPS FOR ORDERING INVESTIGATIONS

 Order X-KUB and Ultrasound in all patients of suspected renal stones (90% of renal stones are



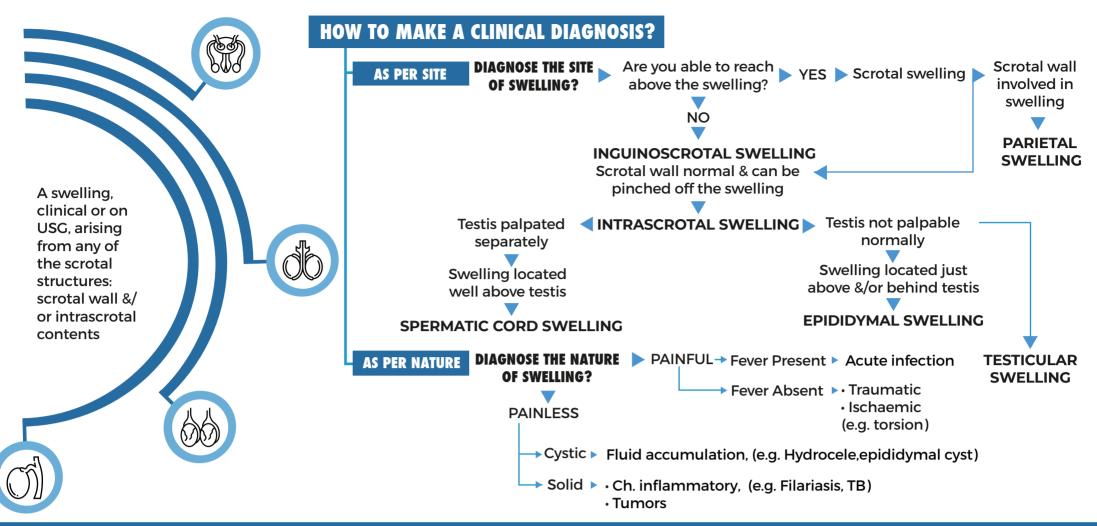
KEEP A HIGH THRESHOLD FOR INVASIVE PROCEDURES



NDIAN COUNCIL OF MENDELT. NEW DELT.

Standard Treatment Workflow (STW) for the Management of SCROTAL SWELLING

ICD-10-N50.89



MAKE A CINICAL DIAGNOSIS

PARIETAL (SCROTAL WALL) SWELLINGS

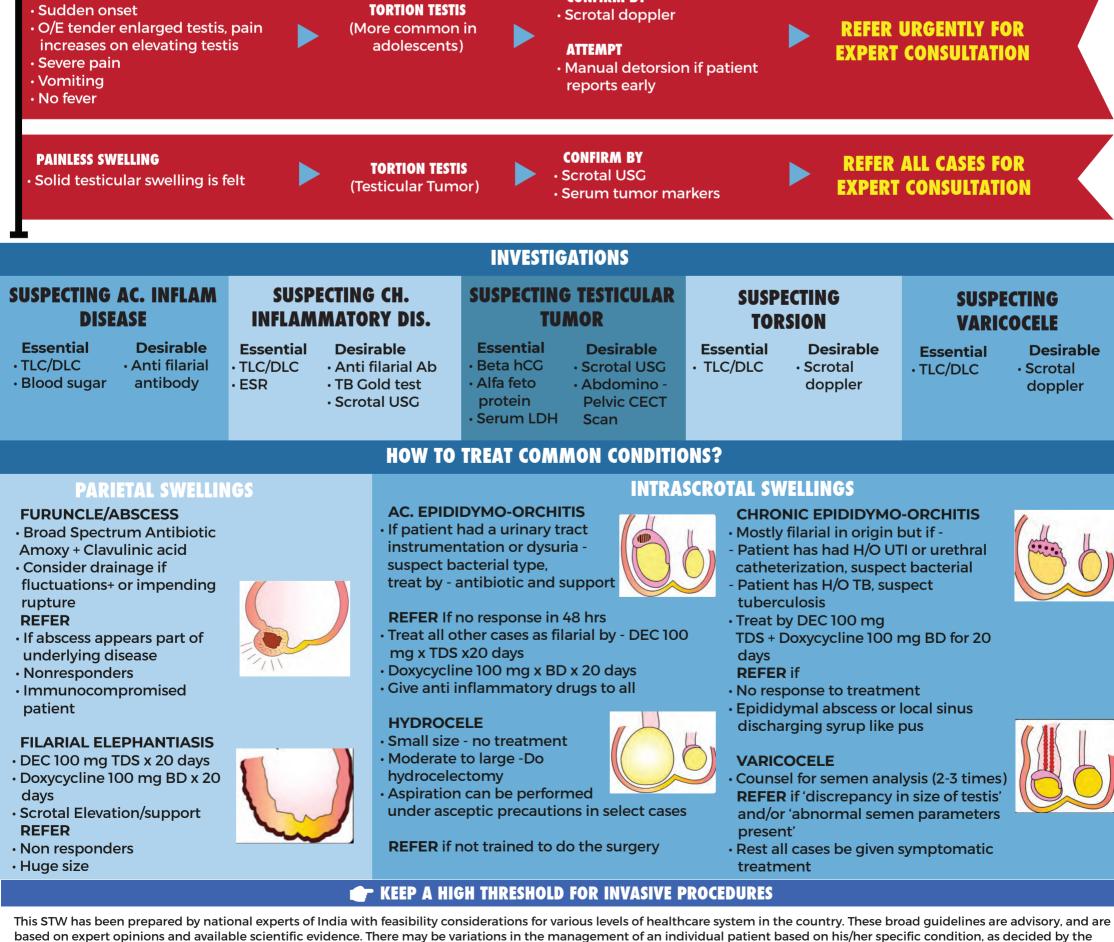
INTRASCROTAL SWELLINGS

	BILATERAL	UNILATERAL		Testicular	Epididymal	Spermatic cord
Ac. Inflammation	• Cellulitis • Fournier gangrene	 Reactionary to epididymo- orchitis Furuncle Abscess 	Cystic	Hydrocele	• Epididymal cyst • Spermatocele	Varicocele
Traumatic	Contusional	Blunt trauma		Deinlass	Painless	
Ch. Inflammation	Filarial Elephantiasis		Solid	PainlessTesticular	 Ch. Filarial epididymitis Ch. Tuberculous Epididymitis Adenomatoid tumor Painful Ac. Epididymitis 	Painless
Fluid Accumulation	 Edema in anasarca, IVC thrombosis Urinary extravasation 	Scrotal wall cysts		tumor Painful		 Lipoma cord Painful Funiculitis
Neoplasm		Melanoma, Scrotal Carcinoma Dermatofibroma;		 Torsion testis Orchitis 		

RED FLAG SIGNS

October/ 2019

CONFIRM BY



treating physician. There will be no indemnity for direct or indirect consequences. Kindly visit our web portal (**stw.icmr.org.in**) for more information.

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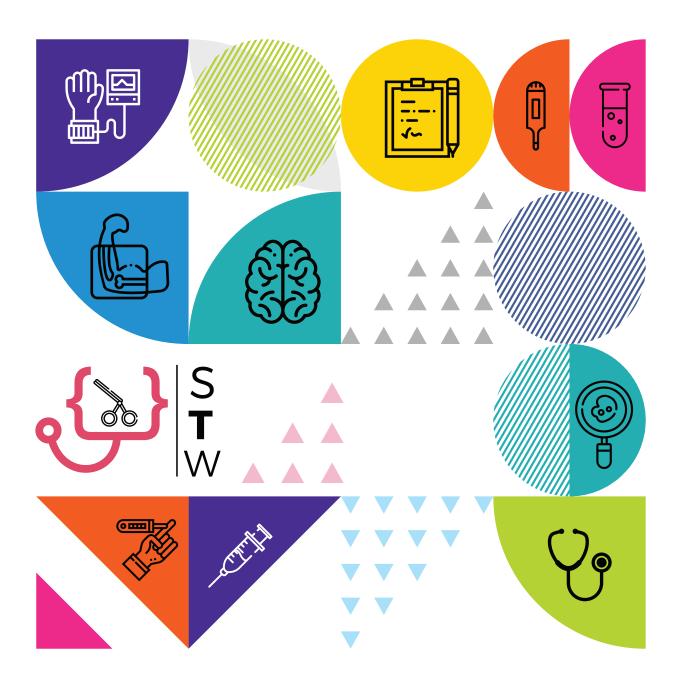








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