

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



Standard Treatment Workflow (STW)

VITILIGO ICD-10-L80

Vitiligo is an acquired sk	Vitiligo is an acquired skin disease characterized by depigmented (white) macules, with a global prevalence of 1-2%						
	NON-SEGMENTAL VITILIGO	SEGMENTAL VITILIGO					
GENERALIZED • Lesions in a g distribution, u affecting trun extremities an • No predilection specific site; a vitiligo vulgar	generalized usuallyAffects the distal extremities and/or face/genitalsnk,face/genitalsnd face on for any also calledtreatment	• Mucosal	 Unilateral with a midline demarcati Onset in childhood Leucotrichia both within and beyond the lesion Usually stabilizes within a year after an initial period of progression Response to medical treatment is variable and most patients may req surgical treatment 				
Generalized vitiligo	Progressive vitiligo with Koebner's phenomenon						
Generunzeu annigo	Koebner's phenomenon	Acrofacial vitiligo	Universal vitiligo Seg	mental vitiligo			
GENERAL PRINCIPLES OF MANAGEMENT							

• Diagnosis is clinical

- Educate patient about the disease
- Assess the psychosocial impact of vitiligo and counsel about the variable/ unpredictable course of disease & expected response to treatment
- In pregnancy, prefer only topical corticosteroids

Decide the treatment plan based on

A Disease activity

- Progressive: new lesions, or spread of existing lesions
- Rapidly progressive: >5 new lesions in last 1 month, or >15 lesions in last 3 months
- Slowly progressive: <5 new lesions in last 1 month, or < 15 lesions in last 3 months
- Stable: no new lesions, no spread of existing lesions
- B Extent of involvement: limited (≤5%) or extensive (>5%)

Limited stable/slowly progressive vitiligo:

Topical treatment- Mid-potent/potent corticosteroids, tacrolimus, topical PUVA/PUVAsol (Avoid prolonged use)

COMMON DIFFERENTIAL DIAGNOSES

Leprosy

- Hypopigmented, not depigmented macules
- Overlying sensory loss
- Enlarged peripheral nerves

· Pityriasis alba

- Extensive stable/slowly progressive vitiligo:
- → Narrow-band ultraviolet B (NbUVB), oral Psoralen + Ultraviolet A (PUVA)/PUVAsol
- Rapidly progressive vitiligo (limited or extensive):
- → Oral corticosteroids (minipulse) and/or
- → Azathioprine/ Methotrexate

Non-responders:

- Consider combining different modalities if unsatisfactory response with monotherapy
- Consider surgical treatment for stable limited vitiligo/ segmental vitiligo (unresponsive to medical treatment)
 Consider camouflage for poorly responsive vitiligo lesions
- Monitoring of patients on systemic treatment
- Height (children), weight, blood pressure and blood sugar in patients on oral corticosteroids
- Complete Hemogram, Liver Function Test in patients on drugs such as Azathioprine, Methotrexate

IMPORTANT COUNSELLING POINTS

- Not the same as leprosy
- Does not spread by touch
- Not caused by certain foods such as milk, curd, lemon, fish etc
- Treatment is available for vitiligo
- Hypopigmented scaly lesions usually on a child's face

Nevus depigmentosus

- Present since birth or early childhood
- Single hypopigmented macule/ segmental lesion

• Multifactorial, predominantly autoimmune

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REFER TO GENERAL PRINCIPLES OF MANAGEMENT						
Acrofacial vitiligo	Stable	 Primary /secondary Level Face, flexures, genitals: Tacrolimus 0.1% ointment BD Other body sites: Betamethasone valera Mometasone/ Fluticasone/ Fluocinolone cream OD (clobetasol NOT to be used) Refer non-responders to higher center after 3 months 	Tertiary Level • Same as in primary/secondary care • Topical PUVA/PUVAsol • Handheld NbUVB			
	Progressive	Refer to higher center	 Topical PUVA/PUVAsol/ Handheld NbUVB (slowly progressive) Levamisole (slowly progressive) Oral steroid (minipulse) and/or Azathioprine/Methotrexate (rapidly progressive) 			
Generalized vitiligo	Stable	Primary /secondary LevelTertiary Level• Face, flexures, genitals: Tacrolimus 0.1% ointment BD• Same as in primary/secondary care • Oral PUVA/PUVAsol• Other body sites: Betamethasone valerate/ Mometasone/ Fluticasone/ Fluocinolone creation OD (clobetasol propionate NOT to be used)• Whole body NbUVB• Refer non-responders to higher center after 3 months• Whole body NbUVB				
	Progressive	Refer to higher center	 Oral PUVA/PUVAsol/ whole body NbUVB (slowly progressive) Levamisole (slowly progressive) Oral steroid (minipulse) and/or Azathioprine/Methotrexate (rapidly progressive) 			
	Primary /se	condary Level	Tertiary Level			
Universal vitiligo	 Sunscreen/photoprotection Refer to higher center 		 Sunscreen/photoprotection Depigmenting agent like monobenzyl ether of hydroquinone 20% may be considered if patient wishes for complete depigmentation 			
Segmental vitiligo	 Primary /secondary Level Face, flexures, genitals: Tacrolimus 0.1% ointment BD Other body sites: Betamethasone valerate/ Mometasone/ Fluticasone/ Fluocinolone cream OD (clobetasol propionate NOT to be used) Refer non-responders to higher center after 3 months 		epidermal grafting, noncultured epidermal suspension			

WITILIGO CAN BE TREATED. TREATMENT DEPENDS ON EXTENT AND ACTIVITY OF DISEASE

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 the website of DHR for more information: (**stw.icmr.org.in**) for more information. ©Department of Health Research, Ministry of Health & Family Welfare, Government of India.