

8th YEAR

08-10 November 2024

InterContinental Dubai - Festival City
United Arab Emirates



The many faces of Cutaneous Tuberculosis

Dr Willie Visser

Head: Division of Dermatology University of Stellenbosch SOUTH AFRICA



Conflict of Interest



No conflicts of interest for this lecture

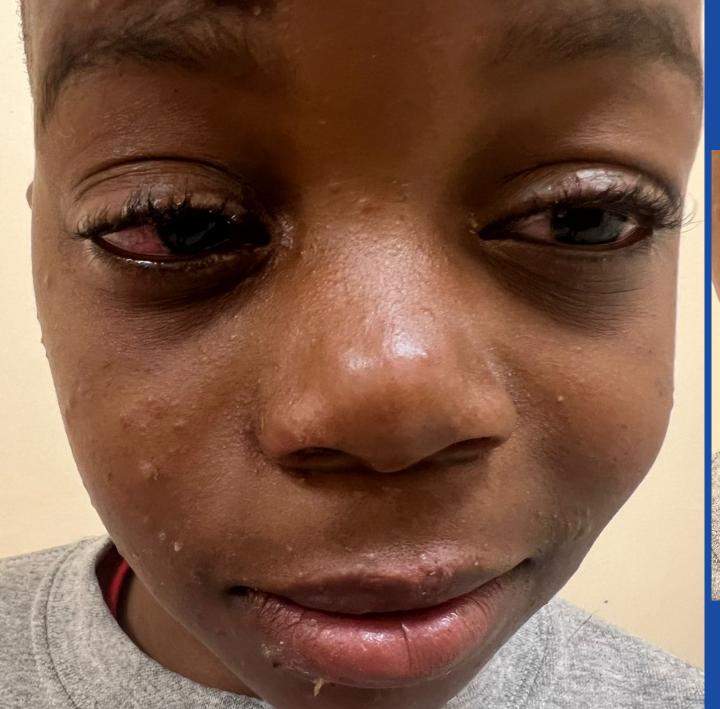


















In this session:



- Classification of cutaneous TB
- Diagnosis of cutaneous TB
- Clinical manifestations of cutaneous TB
- Clinical manifestations of the tuberculids



Classification of cutaneous TB

True cutaneous TB



BEYT ET AL.

TABLE 1. Suggested classification of cutaneous mycobacteriosis and synonymous terms previously used in the medical literature

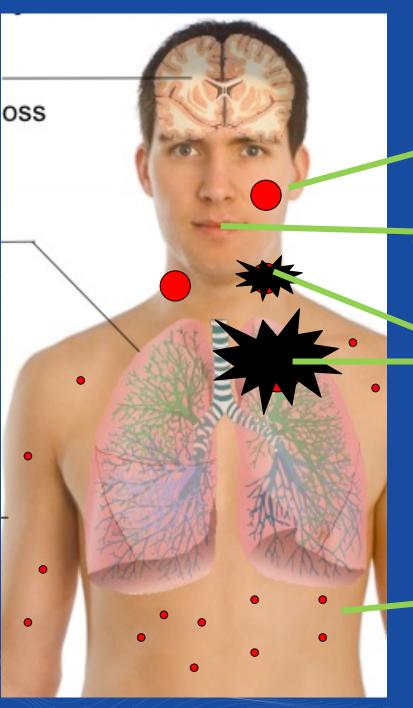
Proposed classification of cutaneous myco- bacteriosis	Terms previously used in literature		
I. Inoculation cutaneous mycobacteriosis	Primary inoculation		
from an exogenous source	Tuberculous chancre		
	Tuberculosis primary complex		
	Tuberculosis verrucosa cutis		
	Warty tuberculosis		
	Verruca necrogenica		
	Prosector's wart		
	Tuberculosis cutis verrucosa		
 Cutaneous mycobacteriosis from an en- dogenous source 			
A. Contiguous spread	Scrofuloderma		
The contract of the contract o	Tuberculosis colliguativa cutis		
B. Autoinoculation	Orificial tuberculosis		
z. ratomountain	Tuberculosis cutis orificialis		
	Tuberculosis ulcerosa cutis et mucosae		
 Cutaneous mycobacteriosis from hematog- enous spread 	a desired esta de		
A. Lupus vulgaris	Lupus vulgaris		
	Tuberculosis luposa cutis		
B. Acute hematogenous dissemination	Acute miliary tuberculosis of the skin		
25 Troute Heminogenesis allocation	Tuberculosis cutis miliaris disseminata		
	Tuberculosis cutis acuta generalisata		
C. Nodules or abscesses	Tuberculous gumma		
C. Troubles of acoccoses	Metastatic tuberculous abscess		











- Lupus vulgaris
- Peri-orifical TB

Scrofuloderma

Miliary tuberculosis



Diagnosis of cutaneous TB

Clinical picture



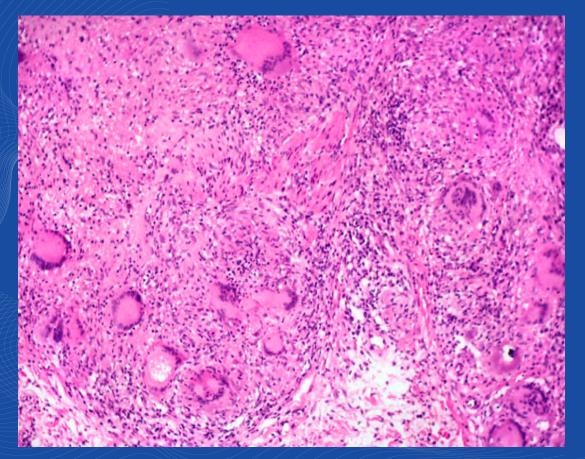
Variable combinations and transitions of papular, nodular, pustular, papulonecrotic, pustulonecrotic, ulcerative, vegetating skin lesions.

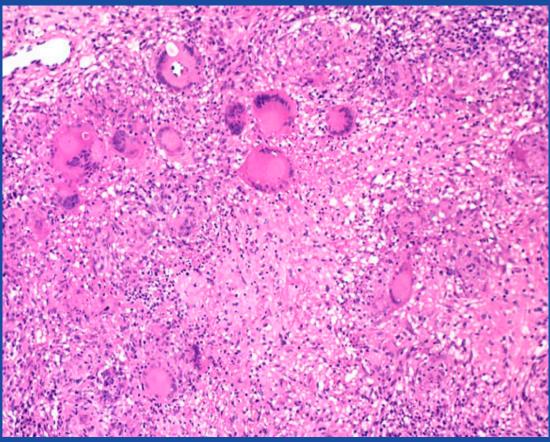






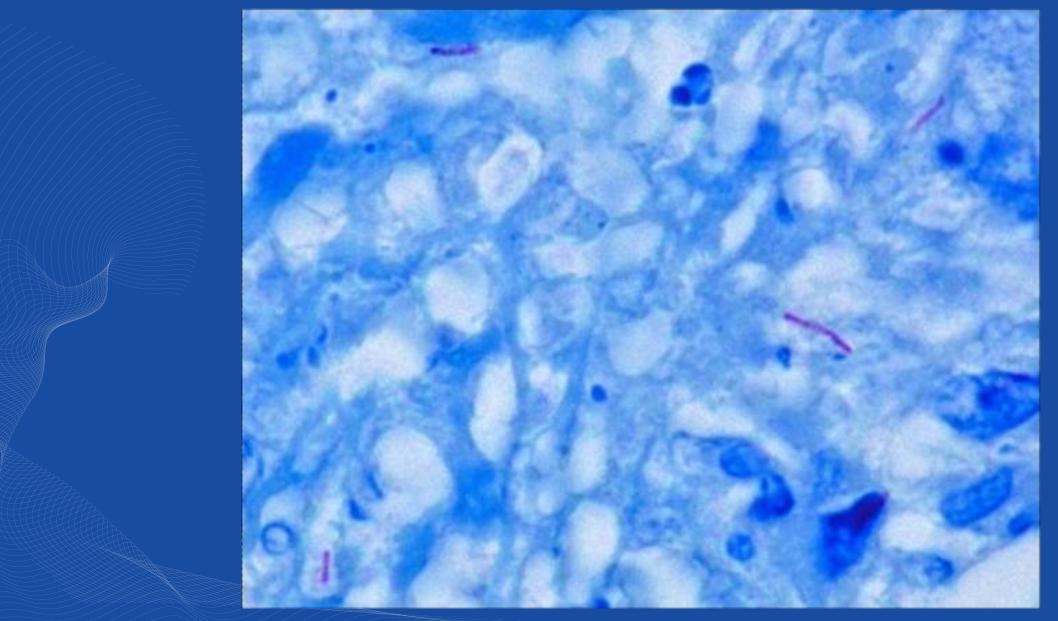
Variable combinations and transitions of **granulomatous inflammation**, mixed acute and chronic inflammatory cells, **necrosis**, **vasculitis**, organisation and fibrosis, other non-specific changes





ZN -stain





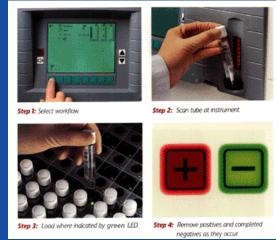


Löwenstein-Jensen

3-8 w



Bactec 2 w



MGIT

Mycobacteria Growth Indicator Tube

1-3 w

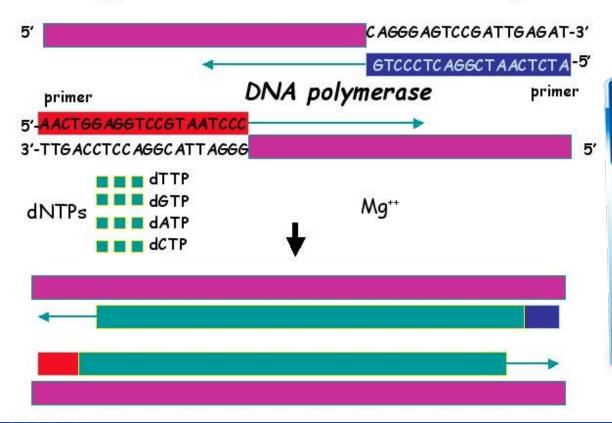
MODS

Microscopic
Observation Drug
Susceptibility
1 w





In PCR 2 non-identical primers are annealed to opposite strands of the DNA template













Response to anti-TB medication



The Adult **Tuberculosis** Guideline

Full prescribing information or Sandoz products included on this brochure is available from your Sandoz Representative

The Paediatric

Tuberculosis

Guideline

Full prescribing information

for Sandoz products included on this

brochure is available from your Sandoz Representative

Regimen 1 (New cases, age above 8 years and adults)

New smear-positive patients, new smear-negative patients and extra-pulmonary TB

Pre-treatment body weight	Two months initial	Four months continuation phase When given SEVEN times a week		
	phase given SEVEN times a week			
	RHZE (150, 75, 400, 275)	RH (150, 75)	RH (300, 150)	
30 - 37 kg	2 tabs	2 tabs		
38 - 54 kg	3 tabs	3 tabs	No. of the last	
55 - 70 kg	4 tabs		2 tabs	
≥71 kg	5 tabs		2 tabs	

Regimen 2 Re-treatment cases)

Pre-treatment body weight			3 rd Month initial phase given SEVEN times a week	Five months continuation phase when given SEVEN times a week			
	RHZE (150, 75, 400, 275)	Streptomycin* (g)	RHZE (150, 75, 400, 275)	RH (150, 75)	E (400)	RH (300, 150)	E (400)
30 - 37 kg	2 tabs	0,5	2 tabs	2 tabs	2 tabs		
38 – 54 kg	3 tabs	0,75	3 tabs	3 tabs	2 tabs		
55 - 70 kg	4 tabs	1,0	4 tabs			2 tabs	3 tabs
≥71 kg	5 tabs	1,0	5 tobs			2 tabs	3 tabs

- * Streptomycin should NOT be given during pregnancy and to those over 65 years.
- ** RH (150, 150) should only be used when treatment is given THREE times weekly in the continuation phase only.
- R Rifampicin, H Isoniazid, Z Pyrazinamide, E Ethambutol



Rimactazid® 150/75 Available in pack sizes of: 56, 84



Rimactazid® 300/150 Available in pack size of:



Available in pack sizes of: 28, 56, 84, 100, 112, 500

& SANDOZ

Marketed by Sandoz SA (Pty) Ltd¹
72 Steel Road, Sparton, Kempton Park
Tel no (011) 929 9000 Fax no (011) 394 5935

'Sandoz SA (Pty) Ltd, Reg No 1990/001979/07



Regimen 3 (Children with tuberculosis - up to the age of 8 years)

Pre-treatment body weight	Two months initial	Four months continuation phase When given SEVEN times a week		
	phase treatment given SEVEN times per week			
	RHZ (60, 30, 150)	RH (60, 30)		
3 - 4 kg	% tablet	% tablet		
5 - 7 kg	1 tablet	1 tablet		
8 - 9 kg	1 ½ tablets	1 ½ tablets		
10 - 14 kg	2 tablets	2 tablets		
15 - 19 kg	3 tablets	3 tablets		
20 - 24 kg	4 tablets	4 tablets		
25 - 29 kg	5 tablets	5 tablets		
30 - 35 kg	6 tablets	6 tablets		

All children with severe forms of tuberculosis (meningitis, spine, peritonitis, miliary, bones) should

Active case-finding is necessary for all children under the age of five years who are in close contact with an infectious TB case. These children should be examined and if found to be healthy, should be given prophylaxis. The recommended regimen is Isoniazid 5 mg/kg daily for six months. Those children found to have tuberculosis, should be treated with a full course of TB treatment.

For more information on management of tuberculosis, refer to the National Tuberculosis Control Programme Guidelines.

Gillicanosis 4 DC Tablain Ray no 23/202 3/0016 Each habite content 150 ng Bisspoor, 75 ng hamistod, 400 ng Pyrazinosisko, 275 ng Bhambotd HCI Billicencandil 3000/150 bilain Ray na 15/202 3/0016 Each habite content 150 ng Bisspoor, 15 ng hamistod, 400 ng Pyrazinosisko, 275 ng Bhambotd HCI Billicencandil 3000/150 bilain Ray na 15/202 3/006 Each habite contents 000 ng Bisspoor, 150 ng benazid. Billicencandil 3000/150 bilain Ray na 15/202 3/006 Each habite contents 000 ng Bisspoor, 100 ng benazid. 150 ng Pyrazinosisko. Billicencandil 7 hed 60/100 bilain Ray na 25/202 3/006 Each habite contents 00 ng Bisspoor, 100 ng benazid. 150 ng benazid.



Rimcure® Paed 3 - FDC Available in pack sizes of: 28, 56, 84



Rimactazid® Paed 60/30 Available in pack sizes of: 28, 56, 84









TUBERCULOSIS

A COMPREHENSIVE CLINICAL REFERENCE

ASSOCIATE EDITORS

John M. Grange • Peter R. Donald • Madhukar Pai Mario Raviglione • Jeffrey R Starke • Wing Wai Yew







H. Simon Schaaf
Alimuddin I. Zumla

FOREWORD BY

Archbishop Emeritus Desmond Tutu



This edition is for sale in Bangladesh, Bhutan, India, Maldives, Nepal, Pakistan and Sri Lanka. This edition is also for sale in selected countries in Africa, namely, Gambia, Ghana, Malawi, Mauritius, Nigeria, Sierra Leone, Angola, Democratic Republic of Congo, Botswana, Swaziland, Lesotho, Zambia, Zimbabwe, Ethiopia, Kenya, Rwanda, Burundi, Tanzania, and Uganda. It may also be sold in Egypt, Sudan, Russia and CIS countries by arrangement with the publisher.

TUBERCULOSIS

A COMPREHENSIVE CLINICAL REFERENCE

EDITED BY

SAUNDERS

H. Simon Schaaf Alimuddin Zumla

ASSOCIATE EDITORS

John M. Grange · Peter R. Donald · Madhukar Pai Mario Raviglione · Jeffrey R Starke · Wing Wai Yew

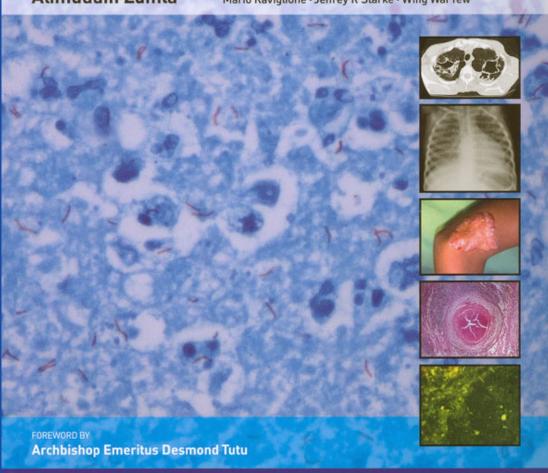


TABLE 2. A PRACTICAL GUIDE TO THE DIAGNOSIS AND MANAGEMENT CUTANEOUS **TUBERCULOSIS**

	Definite cutaneous TB	Probable cutaneous TB	Possible cutaneous TB		
Clinical morphology	Variable combinations and transitions of papular, nodular, pustular, papulonecrotic, pustulonecrotic, ulcerative, vegetating skin lesions.				
Histopathology	Variable combinations and transitions of granulomatous inflammation, mixed acute and chronic inflammatory cells, necrosis, vasculitis, organisation and fibrosis, other non-specific changes				
ZN and / or culture and / or PCR on skin lesion biopsy	+^	-	-		
ZN and / or culture and / or PCR on specimen from origin other than skin	NEFD	+*	2		
Mantoux test result	NEFD	+* (only in children <5 years old)	-		
X-ray findings compatible with TB e.g. lung, bone, etc.	NEFD	+*	ā.		
Response to antiTB treatment	NEFD	NEFD	EFD		

Tuberculosis TB

ZN Ziehl-Neelsen stain

PCR Polymerase chain reaction Not essential for diagnosis NEFD

EFD

Essential for diagnosis (but not necessarily proof of tuberculosis)
When only ZN positive, distinction from environmental (non-tuberculous) mycobacteria is necessary by culture and / or PCR
Any positive establishes the diagnosis of probable tuberculosis





TABLE 2. A PRACTICAL GUIDE TO THE DIAGNOSIS AND MANAGEMENT CUTANEOUS **TUBERCULOSIS**

	Definite cutaneous TB	Probable cutaneous TB	Possible cutaneous TB		
Clinical morphology	Variable combinations and transitions of papular, nodular, pustular, papulonecrotic, pustulonecrotic, ulcerative, vegetating skin lesions.				
Histopathology	Variable combinations and transitions of granulomatous inflammation, mixed acute and chronic inflammatory cells, necrosis, vasculitis, organisation and fibrosis, other non-specific changes				
ZN and / or culture and / or PCR on skin lesion biopsy	+^	-	-		
ZN and / or culture and / or PCR on specimen from origin other than skin	NEFD	+*	¥		
Mantoux test result	NEFD	+* (only in children <5 years old)	-		
X-ray findings compatible with TB e.g. lung, bone, etc.	NEFD	+*	-		
Response to antiTB treatment	NEFD	NEFD	EFD		

TB Tuberculosis ZN Ziehl-Neelsen stain

PCR Polymerase chain reaction Not essential for diagnosis NEFD

Essential for diagnosis (but not necessarily proof of tuberculosis)
When only ZN positive, distinction from environmental (non-tuberculous) mycobacteria is necessary by culture and / or PCR
Any positive establishes the diagnosis of probable tuberculosis





TABLE 2. A PRACTICAL GUIDE TO THE DIAGNOSIS AND MANAGEMENT CUTANEOUS **TUBERCULOSIS**

	Definite cutaneous TB	Probable cutaneous TB	Possible cutaneous TB		
Clinical morphology	Variable combinations and transitions of papular, nodular, pustular, papulonecrotic, pustulonecrotic, ulcerative, vegetating skin lesions.				
Histopathology	Variable combinations and transitions of granulomatous inflammation, mixed acute and chronic inflammatory cells, necrosis, vasculitis, organisation and fibrosis, other non-specific changes				
ZN and / or culture and / or PCR on skin lesion biopsy	+^	-	-		
ZN and / or culture and / or PCR on specimen from origin other than skin	NEFD	+*	F		
Mantoux test result	NEFD	+* (only in children <5 years old)	-		
X-ray findings compatible with TB e.g. lung, bone, etc.	NEFD	+*			
Response to antiTB treatment	NEFD	NEFD	EFD		

TB Tuberculosis

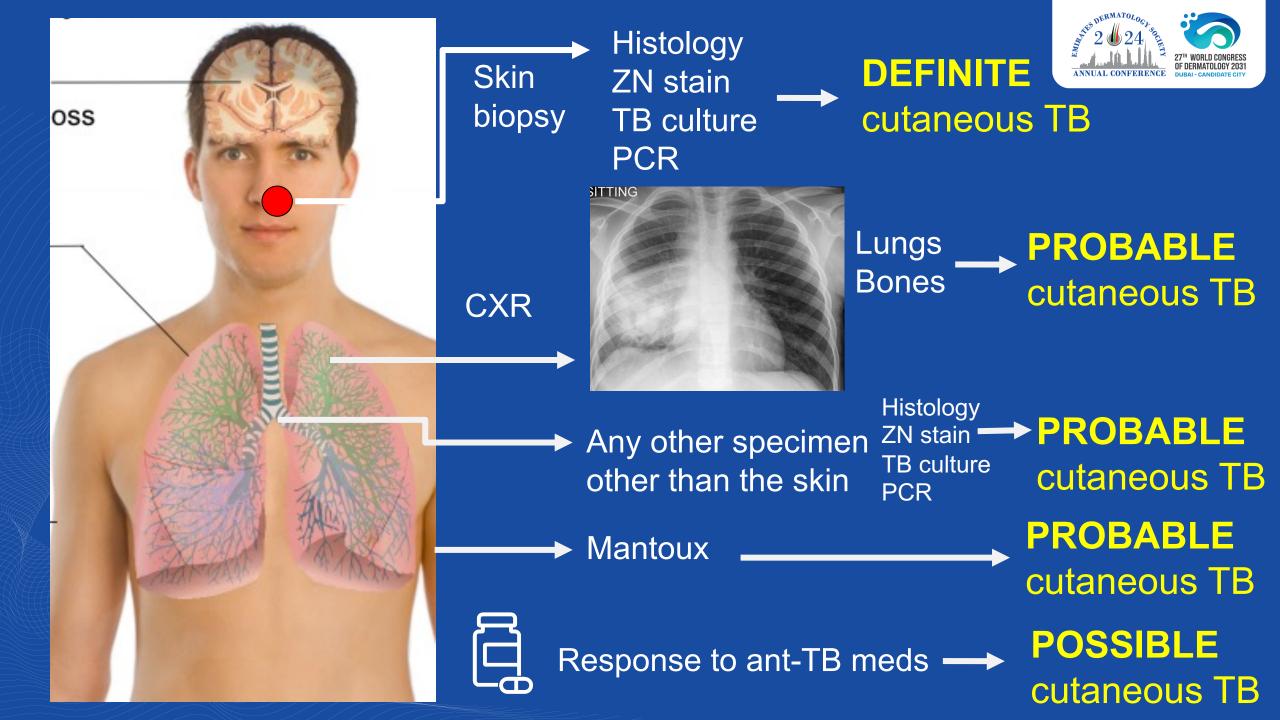
ZN Ziehl-Neelsen stain

PCR Polymerase chain reaction Not essential for diagnosis NEFD

Essential for diagnosis (but not necessarily proof of tuberculosis)
When only ZN positive, distinction from environmental (non-tuberculous) mycobacteria is necessary by culture and / or PCR
Any positive establishes the diagnosis of probable tuberculosis









Examples of cutaneous TB



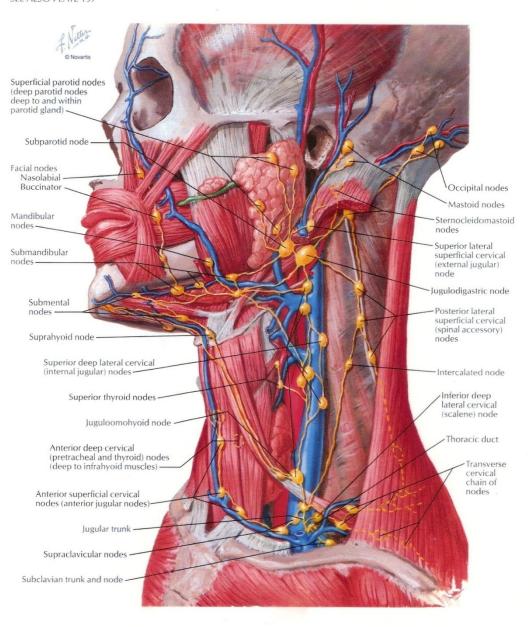






Lymph Vessels and Nodes of Head and Neck

SEE ALSO PLATE 197





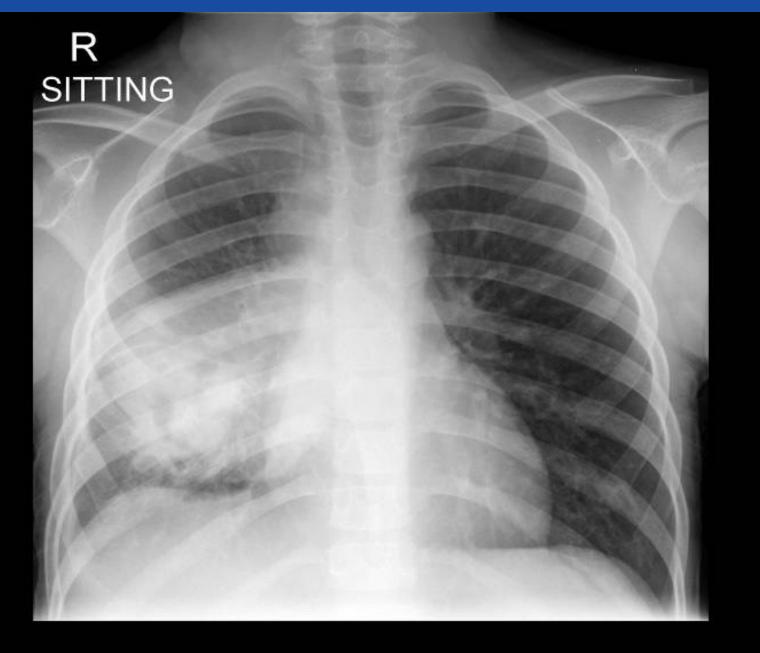




























































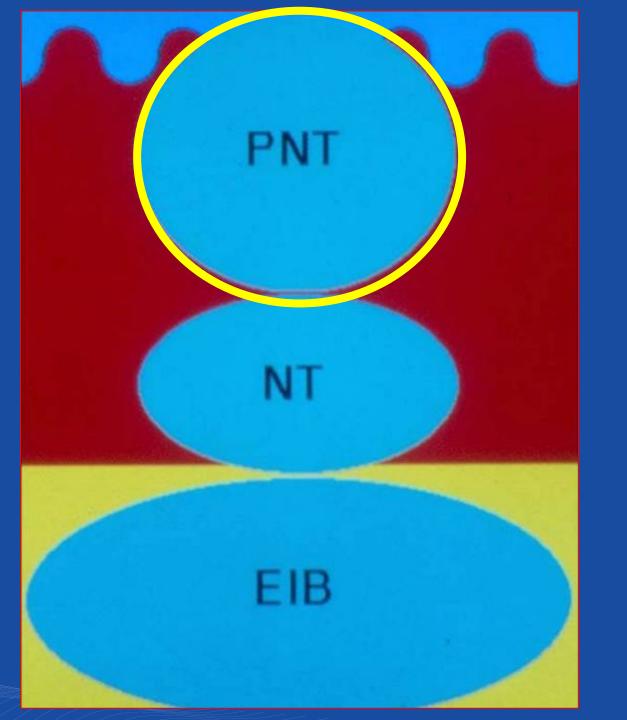






Tuberculids

- Papulonecrotic tuberculid (PNT)
- Erythema induratum of Bazin (EIB)
- Nodular tuberculid (NT)
- Lichen scrofulosorum (LS)
- Phlebitic tuberculid (PT)

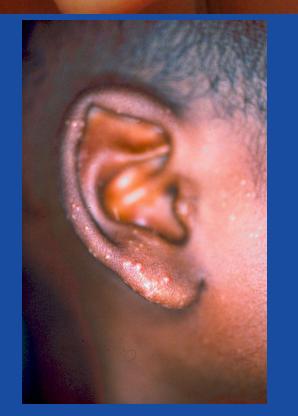
























































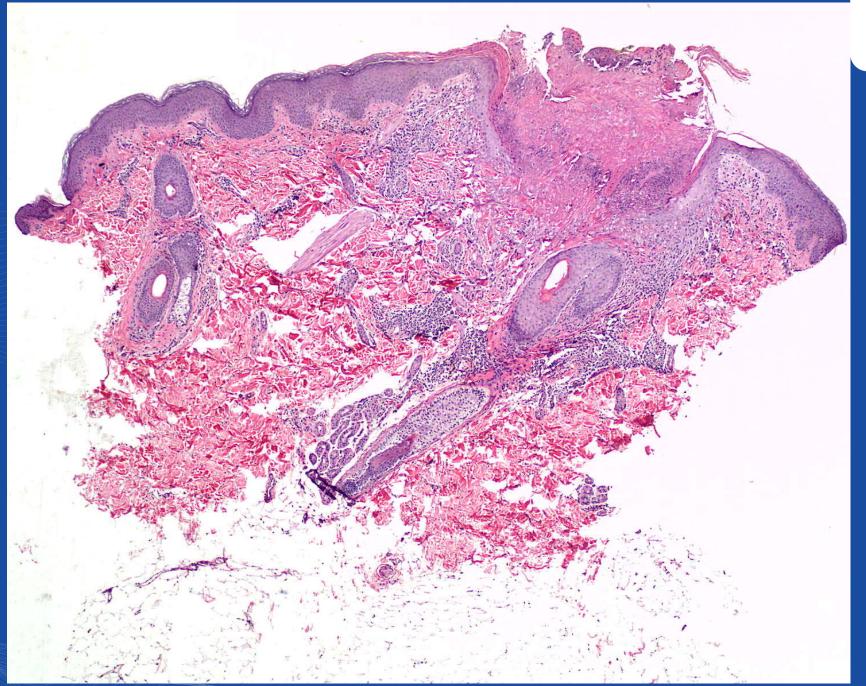






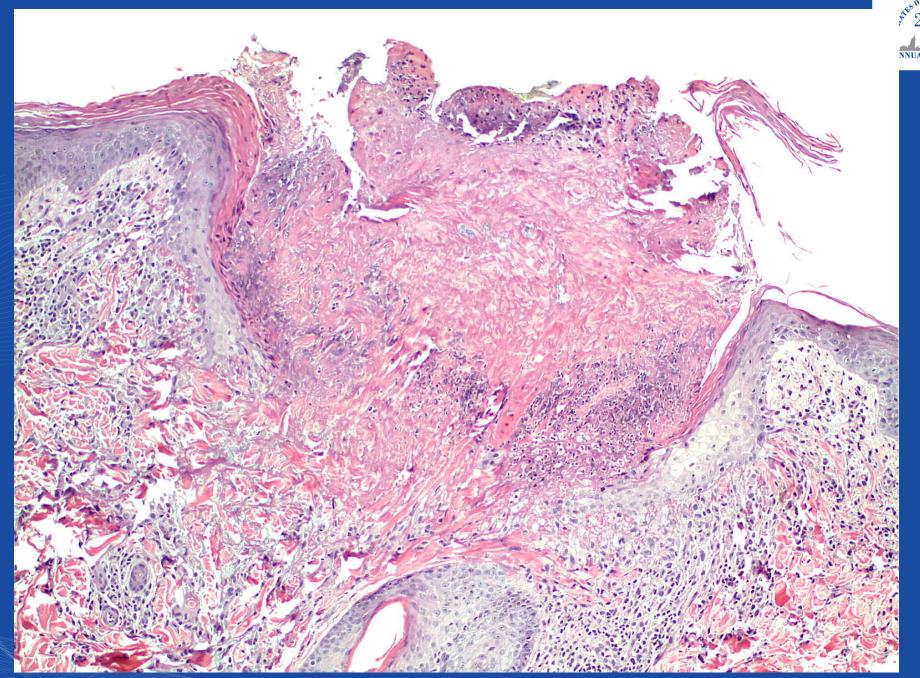




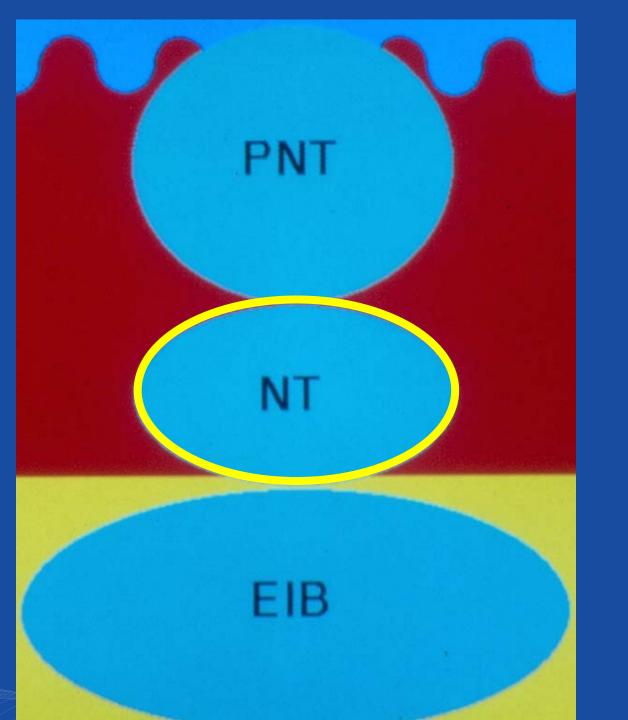


























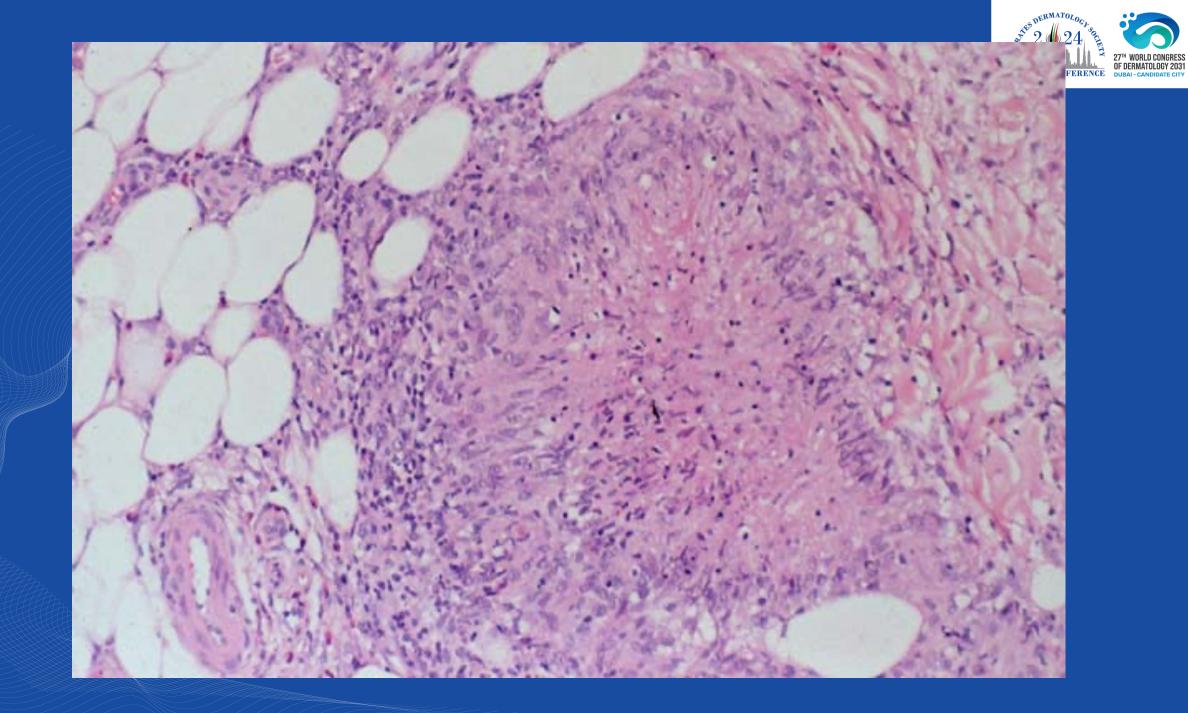


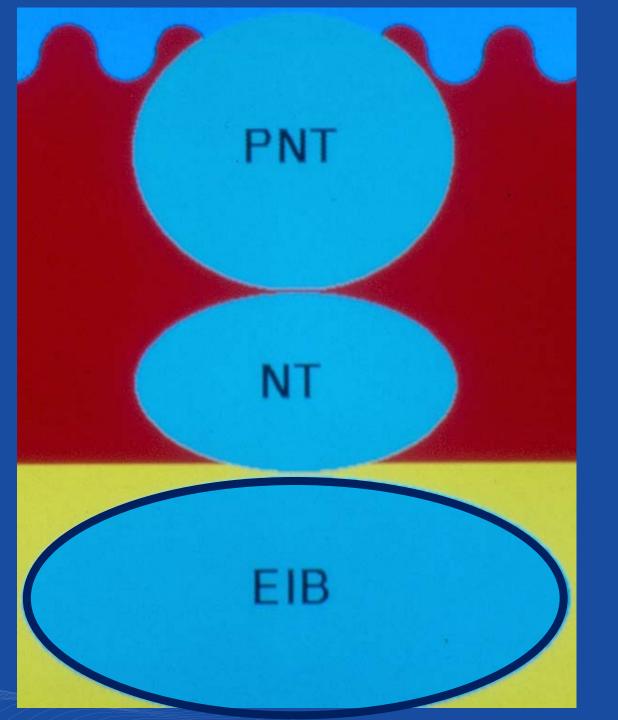
















27TH WORLD CONGRESS OF DERMATOLOGY 2031 DUBAI - CANDIDATE CITY







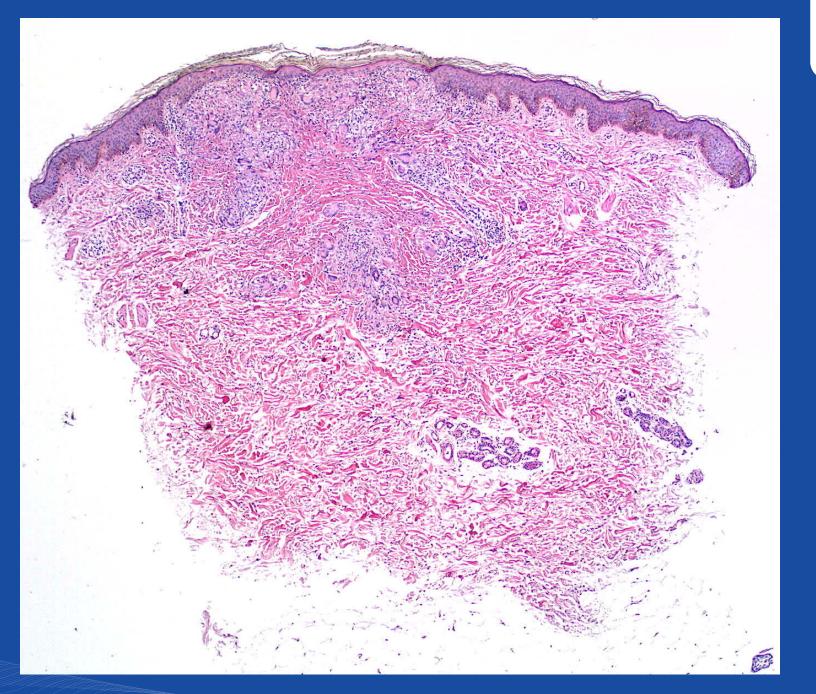






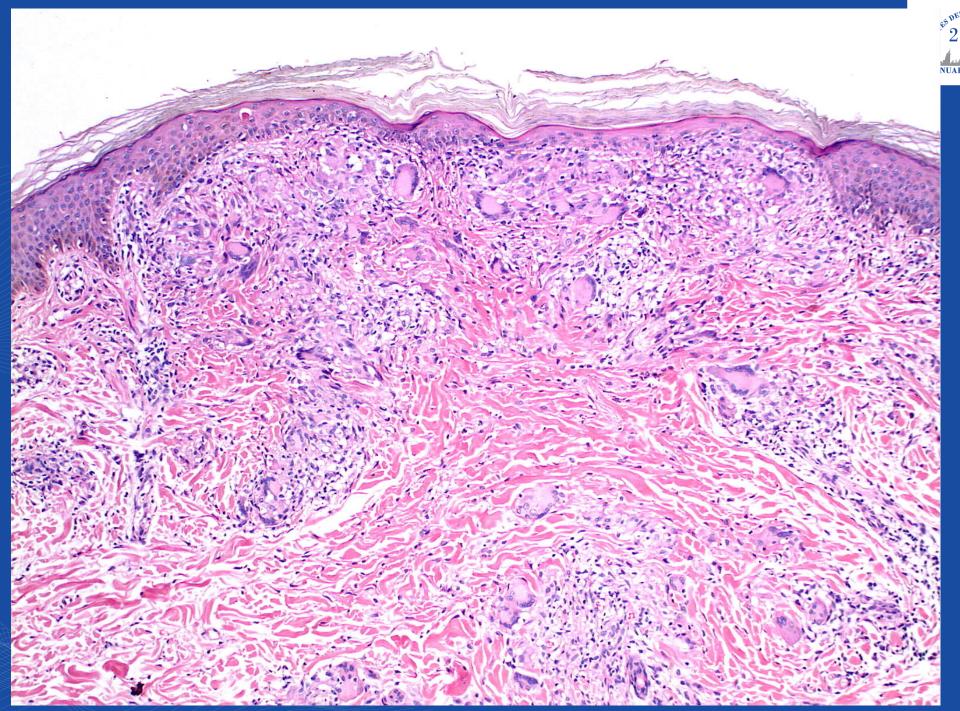














Case report

Superficial thrombophlebitic tuberculide

Hendrick M. Motswaledi, FCDerm (SA), and E. Joy Schulz, MMed (Derm)

From the Department of Dermatology, University of Limpopo and Division of Dermatology, Department of Medicine, University of the Witwatersrand, South Africa

Correspondence

Hendrick M. Motswaledi, FCDerm (SA)
Department of Dermatology
Medunsa Campus
University of Limpopo
South Africa
E-mail: griet@webmail.co.za

Abstract

Background Tuberculides are the result of immunologic reactions to hematogenously spread antigenic components of *Mycobacterium tuberculosis*. There are three recognized tuberculides – papulonecrotic tuberculide, erythema induratum of Bazin, and lichen scrofulosorum. In 1997, in Japan, Hara and coworkers reported five patients with what they called "nodular granulomatous phlebitis," which they proposed was a fourth type of tuberculide. We describe a patient who presented with features identical to those reported by Hara *et al.* in order to draw attention to the previous report and to support the concept of a fourth tuberculide which clinically resembles superficial thrombophlebitis.

Methods A black South African man presented with cord-like thickening of superficial veins on the antero-medial aspects of the lower legs. Nodular swellings were palpable along the course of these veins. There was no evidence of tuberculosis elsewhere in the body, but the patient had a strongly positive tuberculin reaction. Skin biopsies were performed for histologic examination, culture, and polymerase chain reaction (PCR).

Results Histologic examination showed a granulomatous infiltrate localized to the veins in the subcutaneous fat. Stains for acid-fast bacilli and culture were negative, but PCR was positive for *M. tuberculosis* DNA. The lesions responded promptly to antituberculous therapy. **Conclusions** Our patient showed features identical to those of cases described by Hara and coworkers and assigned as a fourth type of tuberculide. As the lesions clinically resemble superficial thrombophlebitis, we propose the term "superficial thrombophlebitic tuberculide" rather than "nodular granulomatous phlebitis."





Int J Dermatol 2006; 45: 1337-1340









Phlyctenular

conjunctivitis/keratoconjuntivitis
= a hypersensitivity reaction to antigens

Staphylococcal Chlamydia

TΒ

Cooccidiodes imitis
Some parasites
Candida
Idiopathic



Tuberculids

Papulonecrotic tuberculid Erythema induratum of Bazin Nodular tuberculid Lichen scrofulosorum Phlebitic tuberculid

Management



- The current consensus: **standard 6-month TB treatment regimen (**2-month intensive phase with rifampicin, isoniazid, pyrazinamide, and ethambutol (RHZE) followed by 4 months of rifampicin and isoniazid)
- **Trial of treatment**: standard TB treatment for 5-6 weeks, then review for response
- Treatment of tuberculids: the same as for other forms of cutaneous TB
- Recommended treatment for Erythema induratum is the standard 6-month regimen, with the addition of a longer treatment period of isoniazid 400mg/day for up to 2 years (11,21,28).
- Pyridoxine should be added in all cases to prevent peripheral neuropathy.

Cutaneous Tuberculosis: A Retrospective Review at a South African



Tertiary Dermatology Unit

Principal investigator

Dr Barbara van der Westhuizen Division of Dermatology, Stellenbosch University



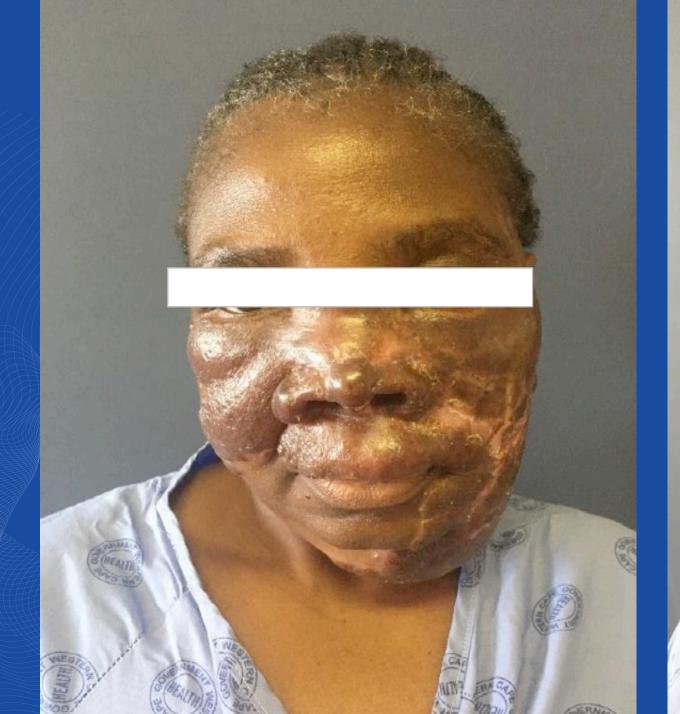
Study period of **5 years**, from 1 January 2018 to 31 December 2022



Variable	Frequency, n	Percentage, %
Age		
<10y	8	27.6
11-20y	2	6.9
21-30y	5	17.2
31-40y	7	24.1
41-50y	5	17.2
51-60y	2	6.9
Gender		
Male	18	62.1
Female	11	37.9
HIV infection		
Yes	4	13.8
No	23	79.3
Unknown	2	6.9
Known TB contact		
Yes	2	6.9
No	10	34.5
Not documented	17	58.6

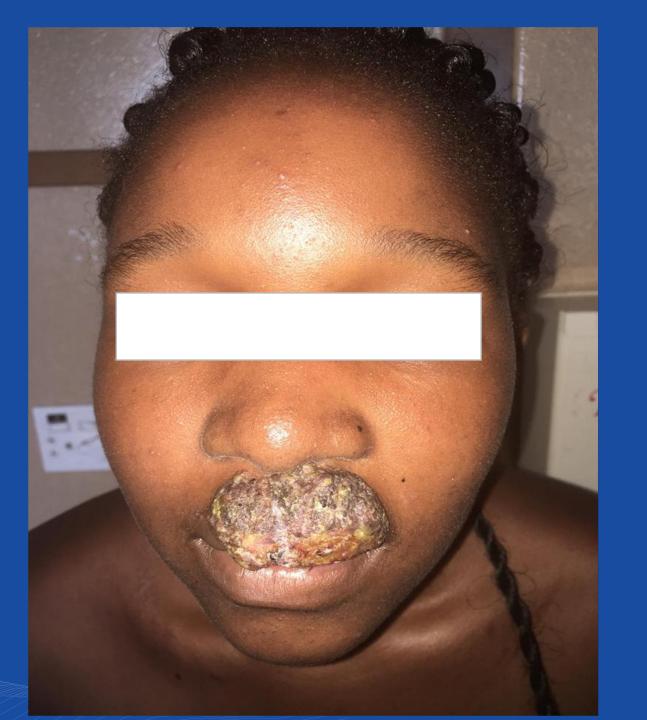


Morphology by subtype			
Erythema induratum of Bazin	Nodules	14	100
Lupus vulgaris	Plaques	5	55.6
	Papules and plaques	2	22.2
	Nodules	1	11.1
	Plaques/nodules	1	11.1
	with ulceration		
Papulonecrotic tuberculid	Papules	3	100
Scrofuloderma	Plaques/nodules	2	100
	with ulceration		
Miliary TB	Papules	1	100

























































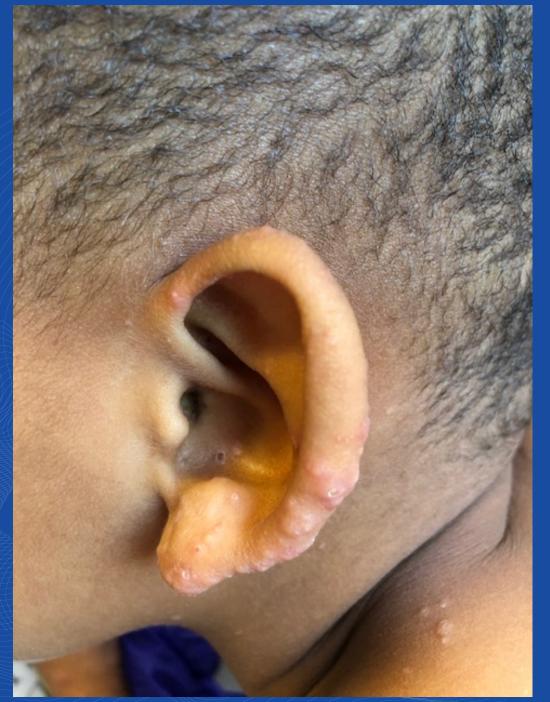




















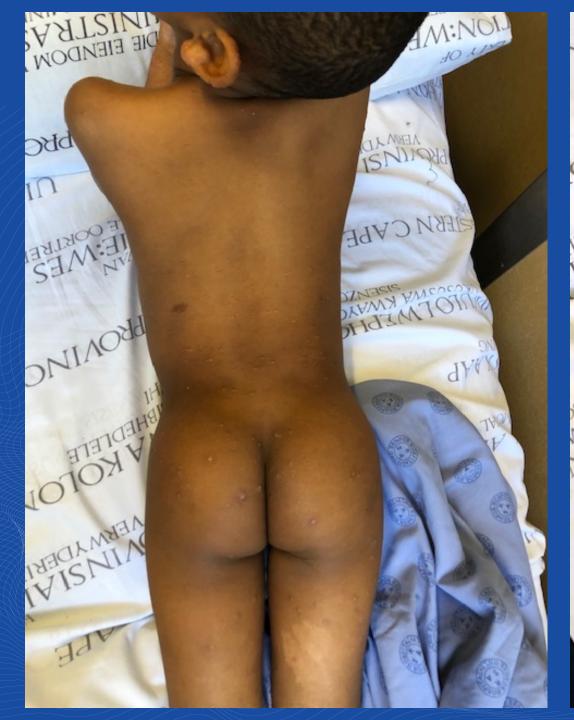












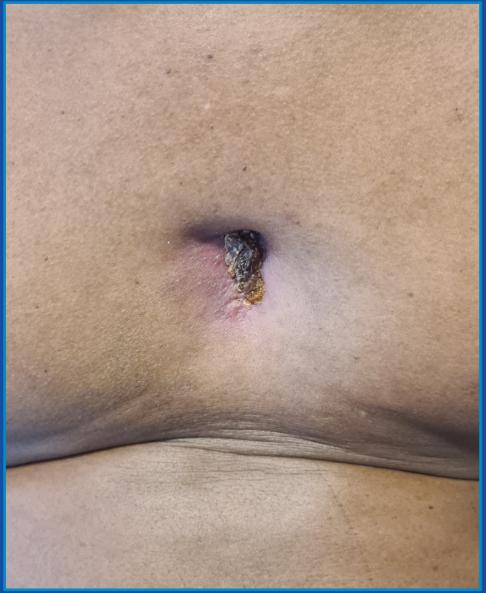


Image 14,15,16. At 4 months









At 6 weeks

Baseline









At 10 weeks













Conclusion



- Cutaneous TB is uncommon
- Cutaneous TB has diverse morphologies
- Diagnosing cutaneous TB is challenging
- Awareness of cutaneous TB could aid in early diagnosis and treatment

