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DOCUMENTO DE CONSENSO

[Translated article] Spanish Academy of Dermatology and Venereology (AEDV) expert recommendations for the management of sexual transmitted parasitosis. Scabies, and pediculosis pubis

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Abstract Sexually transmitted infections are communicable diseases where the pathogen is transmitted through sexual contact. The Sexually Transmitted Infections Working Group of the Spanish Academy of Dermatology and Venereology (AEDV) is engaged in the drafting of documents to guide dermatologists and health care personnel who treat Spanish patients with these infections. This document analyzes the epidemiological, clinical, therapeutic, and control characteristics of 2 sexually transmitted parasitosis: scabies due to *Sarcoptes scabiei* var. *hominis*, and pubic pediculosis due to *Phthirus pubis*. Both parasitoses share a sort of mixed spread through sexual and community transmission regardless of the route through which the infection was initially acquired. This specific feature creates particularities in the management and control of the infestation.

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PALABRAS CLAVE

Escabiosis;
Sarna;
Pediculosis púbica;
Ladillas;
Infecciones de
transmisión sexual;
ITS

Recomendaciones de expertos de la Academia Española de Dermatología (AEDV) para el manejo de las ectoparasitosis de transmisión sexual. Escabiosis y pediculosis púbica

Resumen Las enfermedades transmisibles en las que el patógeno se transmite por contacto sexual se denominan infecciones de transmisión sexual. La Academia Española de Dermatología (AEDV), a través de su grupo de trabajo de infecciones de transmisión sexual, se ha propuesto elaborar documentos de recomendaciones, destinados a la orientación de los dermatólogos y del personal sanitario que atiende en España a personas afectadas de estas infecciones. El presente documento analiza las características epidemiológicas, clínicas, terapéuticas y de control de 2 parasitosis consideradas de transmisión sexual: la escabiosis, causada por *Sarcoptes scabiei* var. *hominis*, y la pediculosis púbica, causada por el *Pthirus pubis*. Ambas parasitosis tienen en común que pueden diseminarse de forma mixta, por vía sexual y comunitaria, independientemente del modo de adquisición de la infestación. Esta peculiaridad genera particularidades en el control de la infestación.

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Introduction

The aim of this document is to establish expert recommendations from the Spanish Academy of Dermatology and Venereology (AEDV) on the management and control of sexually transmitted ectoparasitic infections (STELs), as part of the project of developing AEDV recommendations on the management of sexually transmitted infections (STIs).

The goals are to adapt these recommendations to the scientific evidence currently available and unify criteria and strategies for prevention, detection, diagnosis, treatment, and individual and community control of STELs across different demographic, epidemiological, clinical, and health socio-sanitary situations. The document anticipates a plan for updating every 5 years.

For the effective management of these community-transmitted ectoparasitic infections, a comprehensive approach is required, involving close collaboration among primary care health workforce, dermatologists, infectologists, epidemiologists, and preventive medicine specialists, among others, leading to a coordinated response between them and with those responsible for the surveillance and control policies of communicable diseases. Therefore, this document is not only intended for dermatologists but also for all health care workers who manage scabies and those who establish prevention and control policies.

As [supplementary data](#), annexes are provided with information on scabies, its differential diagnosis ([supplementary data. Annex 1](#)), patient information sheets ([supplementary data. Annex 2](#)), oral ivermectin dosing ([supplementary data. Annex 3](#)), information for workers ([supplementary data. Annex 4](#)), and a checklist ([supplementary data. Annex 5](#)) for affected institutions, frequently asked questions ([supplementary data. Annex 6](#)), video with audiovisual support material for diagnosis, treatment, and control ([supplementary data. Annex 7](#), also available at <https://youtu.be/lP8McE90Cz4>, English version, and <https://aedv.fundacionpielsana.es/wikiderma/escabiosis>, Spanish version), and a table with the levels of evidence and grades of recommendation used ([supplementary data. Annex 8](#)).

Justification

A characteristic of sexually transmitted parasitic infestations such as scabies and pediculosis is their potential for transmission through sexual contact as well as acquisition within the community. Clinical and epidemiological diagnoses are important for individual and sexual contact control and, eventually, community control. Additionally, their presence is suggestive of other possible associated concurrent STIs, including HIV.

Regarding scabies, in recent years, different European countries have reported on a suspected increased incidence and decreased efficacy of treatments indicated in the clinical practice guidelines.¹⁻⁶ Inadequate therapeutic compliance and prevention measures, and decreased efficacy of scabicides, may be contributing to insufficient control.⁷

Pubic lice infestation has not raised alarms of incidence or lack of control, nor has it been demonstrated that it acts as a vector in the transmission of other diseases. However, among affected individuals, an increased incidence of other STIs such as syphilis, gonorrhoea, and chlamydia has been reported.^{8,9}

Methods

The authors reviewed the main international guidelines, including guidelines drafted in Europe,⁹⁻¹¹ Japan,¹² Germany,¹³ the United Kingdom,¹⁴ and the U.S. Centers for Disease Control and Prevention (CDC),^{15,16} and protocols from national and local health services and epidemiological surveillance from Spain, published or provided by dermatologists.¹⁷⁻²⁷ Additionally, in August 2023, relevant scientific publications were consulted in the PubMed, EMBASE, and Cochrane databases, without date restrictions, with available abstracts, and written in English, French, or any of the official languages of Spain. Case descriptions were excluded. The obtained works and their citations were reviewed to identify other relevant studies.

Both the levels of evidence and grades of recommendation were evaluated using the criteria established by the

Oxford Center for Evidence-Based Medicine, version 2009.²⁸ (supplementary data. Annex 8).

Drug prices were consulted in the nomenclature of pharmaceutical products available in pharmacies of the Spanish Ministry of Health as of September 2023.²⁹ The approximate price of drugs available in magistral formulation was calculated by requesting a quote from different pharmacies subject to the regulation of magistral formulation of drugs from the Spanish autonomous communities of Andalusia, Cantabria, Castilla-La Mancha, Catalonia, and Madrid.

External review

Two experts, one representing the International Alliance for the Control of Scabies (IACS) and the other from the World Health Organization (WHO), both Spanish speakers, reviewed the document. Their recommendations were carefully analyzed and considered.

Scabies

What is scabies?

Human scabies, is a communicable disease resulting from infestation of human skin by a mite of the family *Sarcoptidae*. References to its existence date back to ancient times.^{30,31}

Etiology

The causative agent of scabies is *Sarcoptes scabiei* var. *hominis*, a 0.30 mm x 0.50 mm arthropod (size of adult females) at the limit of human eye visibility.³² Its complete life cycle, which extends from 9 to 15 days, occurs in humans, who serve as its reservoir. Mated females burrow into the epidermis, where they lay their eggs. Female survival and infectivity in the environment are higher when relative humidity is high, and temperature drops below 20 °C.³³ Under normal conditions, the mite can survive for 3 days outside the host.^{32,34,35}

Transmission

Transmission occurs through close physical contact for, at least, 20 minutes, with another infected individual, typically a sexual partner or a household member.³⁶ Cases of crusted scabies (see “Clinical forms” section below) harbor a high parasite load (hyperinfestation), where brief contacts may be sufficient for transmission.^{14,37}

The contagious period starts when the parasite is acquired. Transmission via fomites is unlikely,^{38,39} except for cases of hyperinfestation, where it poses a real risk.^{14,37,40}

Transitory self-resolving infestations have been reported, but never cross-infestation between humans and other animals that complete the parasite’s life cycle.^{41–44}

Pathophysiology

Pruritus, a key symptom, and skin lesions are the result of the host’s immune response to the mite, its excretions, or

eggs and may persist for weeks after parasitological cure. The time elapsed between the initial infestation and this reaction ranges between 2 and 6 weeks, which facilitates spread from asymptomatic carriers.^{38,45}

An episode of scabies does not confer immunity against future infestations. Symptoms often appear 1 to 3 days after exposure in infected individuals with previous episodes of scabies.

Epidemiology

Scabies has a worldwide distribution, with highly variable prevalence across different regions. Scabies has an estimated prevalence of nearly 200 million people each year, and an incidence of 455 million cases per year, which represents a significant global disease burden.⁴⁶ Scabies has been categorized as a “neglected tropical disease” by the WHO,⁴⁷ which prioritizes its control in its 2021-2030 roadmap.⁴⁸

Cyclical patterns and seasonal variations of prevalence are described, but no predictive models have ever been confirmed.^{46,49,50} Situations fostering vulnerability, such as migratory movements, can lead to serious situations of very high prevalence.⁵¹

Scabies is not a notifiable disease, with the exception of outbreaks (see the “Situation Diagnosis” section below), which in Spain and other countries, require reporting. This poses a challenge in accurately determining the prevalence and trends of incidence curves.^{1,52,53}

As it happens in other countries, Spain has seen growing reports of rising case numbers and acaricide use in recent years.^{6,54}

Clinical presentation

Clinical forms

Diagnosis is based on history-taking and recognition of the clinical signs associated with scabies.

The cardinal symptom is intense pruritus, typically sparing the head and face and worsening in the evening hours.

The most specific clinical sign is the burrow—visible to the naked eye—a 2 mm x 15 mm thin, sinuous linear trajectory. Although it is the pathognomonic lesion, it is often hidden by excoriation or superinfection. Other types of lesions, such as papules (figure 1a) and genital nodules in cases of sexual transmission (figure 1c), are more common.

Clinical presentation shares signs and symptoms with other dermatopathies and manifests differently in infants and individuals with a deficient immune response to the mite.^{55,56} Various clinical presentations are shown in Table 1. Figures 1a-i illustrate several clinical and microscopic images.

Complications

Bacterial superinfection, favored by scratching, is the most common complication. The most widely reported bacteria involved in the infection are *Staphylococcus aureus* and *Streptococcus pyogenes*, and the usual clinical presentation is impetigo (figure 1f). In untreated or weakened patients, it can progress into deep soft tissue infections such as cellulitis and sepsis.^{55,57}

Table 1 Clinical forms of scabies.

	Frequency/Characteristics	Pruritus	Preferred location	Lesions
Classic scabies	Common form	<ul style="list-style-type: none"> - Intense - Nocturnal predominance 	<ul style="list-style-type: none"> - Interdigital spaces - Wrists - Axillae - Waist - Feet - Buttocks - Thighs (inner surface) - Areola (females) - Genitals (more common in males) 	<ul style="list-style-type: none"> - Burrows - Papules - Erosions - Vesicles - Pustules - Eczema/lichenification - Nodules (common in genitals) - Blisters (figure 1g)
Infantile acropustulosis	Common form in children < 1 year	<ul style="list-style-type: none"> - Often manifests as restlessness or crying - Scratching may be absent - Mild or absent 	<ul style="list-style-type: none"> - Palms - Soles - Disseminated, including face and head 	<ul style="list-style-type: none"> - Vesiculopustules (figure 1d) - Nodules - Very prominent burrows
Crusted scabies (formerly known as "Norwegian" scabies) (hyperinfestation)	<ul style="list-style-type: none"> - Very rare - Immunocompromised, elderly, malnourished - Large number of viable mites (high contagiousness) - Increased risk of complications, such as severe infections 		<ul style="list-style-type: none"> - Broad cutaneous surfaces - Unusual locations in the classic form (palms, soles, head, face, and nails) (figure 1b) 	<ul style="list-style-type: none"> - Scaling - Crusts - Hyperkeratosis - Erythroderma - Thickening or irregularities in nails - Rarely, burrows
Atypical scabies	<ul style="list-style-type: none"> - Rare - Atypical signs and symptoms, which complicates diagnosis - Mimics known dermatoses (supplementary data. Annex 1) 	Variable	Variable	- Variable (figure 1e)



Figure 1 Clinical and microscopic images of scabies; a) classical form in a young adult with skin type 3 exhibiting the characteristic burrows (arrows) and papules (arrowheads) evident at typical locations on the hand; b) crusted or Norwegian form (hyperinfestation) in an elderly bedridden woman with skin type 2, nail abnormalities, and discreet crusted lesions on her pinky finger, posing a challenge for diagnosis; c) genital nodules and erosions (arrow) and burrow (arrowhead) in a young man with skin type 5; d) infantile acropustulosis, pustules on the sole of a 3-month-old baby with skin type 6; e) atypical form with disseminated eczematous lesions (abdomen shown) in a middle-aged woman with skin type 3; f) classic impetiginized form, pustules in typical scabies location on the hand of an 8-year-old girl with skin type 6; g) blistering atypical form, with blisters on the side of the hand (arrow) and a burrow on the thumb (arrowhead) in a young woman with skin type 3; h) image of the burrow obtained using a manual dermatoscope. At one end, the dark triangle consistent with the anterior section of the mite can be seen (arrowhead). The sinuous linear path starting here ends in a triangular image (arrows). These structures make up the “delta wing sign”; i) image of the mite (arrow) and its eggs (arrowheads) obtained through optical microscopy of scraping of one of the burrows.

Pruritus prevents rest and, along with the stigma associated with the disease, causes significant psychological distress.⁵⁸

Diagnosis

Diagnostic criteria

The IACS criteria (Table 2), which include typical lesions and symptoms, and epidemiological data, allow for standardized diagnosis with varying levels of rigor. This includes confirmed

(level A), clinical (level B), or suspected scabies (level C). The utility of the IACS criteria is limited in clinical forms other than the classical presentation.⁵⁹

The diagnostic gold standard is the visualization of the mite under an optical microscope. The 10x magnification handheld dermatoscope—a more accessible technique—allows for the visualization of different structures and non-invasive confirmation diagnosis (figure 1i). A brown triangular figure at its distal end is consistent with the anterior section of the mite. This figure (delta), along with

Table 2 The IACS criteria.**Confirmed scabies**

Visualization of, at least, one of the following:

- A1: mite, eggs, or feces on optical microscopy of skin samples
- A2: mite, eggs, or feces using high-resolution imaging modalities
- A3: mite on dermoscopy

Clinical scabies

Presence of, at least, one of the following:

- B1: burrows
- B2: typical lesions in male genital region
- B3: typical lesions with typical distribution and 2 clinical history criteria (H)

Suspected scabies

Presence of one of the following:

- C1: typical lesions with typical distribution and 1 element of clinical history (H)
- C2: atypical lesions or atypical distribution and 2 elements of clinical history (H)

Elements of clinical history

- H1: itching
- H2: contact with an infected individual

IACS, International Alliance for the Control of Scabies.

the triangular image at the opposite end and the sinuous path that connects them (trail), is the “delta wing sign.”⁶⁰ (figure 1h).

Various diagnostic aid techniques have been described⁶¹ (Table 3 and supplementary data. Annex 6).

Situation diagnosis

To determine whether there is an outbreak and establish control measures, it is necessary to know if we are dealing with one or multiple cases, classical or hyperinfestation forms, and recognize if vulnerable groups are affected, such as displaced or institutionalized populations (schools, hospitals, refugee camps, nursing homes, penitentiaries...) ^{12,17,21,62} (figure 2).

Cure diagnosis

Cure is defined by the absence of new lesions and the absence of parasites in 2 different tests performed on the remaining lesions over 2 consecutive weeks.^{12,13}

After parasitological cure, burrows may still be visible, and dermatitis and itching may still go on for weeks. Topical scabicides are irritants and may contribute to this persistence.⁶³

Figure 2 illustrates the diagnostic algorithm of scabies.

Treatment

Therapeutic goals are to eliminate the parasite and its eggs, resolve the symptoms, eradicate the signs and complications associated with the infestation, and control its spread. Patients should receive written information (level of evidence 5, grade of recommendation D)¹⁰ (supplementary data. Annex 2).

There are currently 2 marketed drugs in Spain: topical permethrin and oral ivermectin, both regarded as first-line therapies. Other treatments—available by compounding—include topical sulfur, topical ivermectin, and benzyl benzoate. Drug selection should consider the patient’s preferences and potential contraindications. Not all contacts involved need to use the same scabicide. Since

the mite’s life cycle occurs in the stratum corneum, topical treatment is effective in most cases.

Topical treatments should be applied as described in Table 4, regardless of the infection route or the location of signs and symptoms.

The most relevant information on drugs available in Spain is shown in Table 5 .

Treatment recommendations differ depending on the form of scabies, the patient’s clinical condition, and the epidemiological situation (figure 3).

Drugs in the pipeline, or still not available in Spain

Several macrocyclic lactones are in the pipeline. Moxidectin, which has a good safety profile, has been approved by the U.S. Food and Drug Administration (FDA) to treat onchocerciasis in humans. Compared to ivermectin, moxidectin has greater in vitro scabidical activity, cutaneous bioavailability, and half-life (20 to 40 days, which is longer than the parasite’s life cycle), which would allow for single-dose treatment.^{64,65}

Spinosad, in a 0.9% topical suspension, is an insecticide that has been approved in the United States to treat scabies. It has a high safety profile.⁶⁶

Malathion, crotamiton, and lindane are no longer available in Spain, the latter having been withdrawn due to its neurological toxicity.

Combined therapies

The combination of topical permethrin plus oral ivermectin in a single dose of 200 µg/kg was more effective than monotherapy to treat classical scabies. However, the methodology of the only study in which it has been evaluated does not allow us to determine the level of evidence or grade of recommendation of this combination.⁶⁷

For hyperinfestation, a combination of topical and oral scabicides is advised. Topical treatment is repeated daily for a week and then every 48 hours until definitive cure. Oral ivermectin is administered on days 1, 2, and 8 in mild cases and on days 1, 2, 8, 9, 15, 22, and 29 or until a definitive cure

Table 3 Diagnostic aid and confirmation techniques.

	Technique	Facilitates viewing the burrow	Allows visualization of the mite	Non-invasive technique	In vivo visualization of the mite	Accessibility/ Price availability	Learning difficulty
Transillumination	Light source applied to the palmar surface of the interdigital fold	✓	✗	✓	✗	↑↑	↓ ↓
Ink stain	Apply ink to the burrow and clean the excess	✓	✗	✓	✗	↑↑	↓ ↓
UV light	Illuminate with LED 365 nm ^{61,68}	✓	✗	✓	✗	+/-	↓ ↓
Dermatoscope	Manual device with 10x magnification and illumination	✓	✓	✓	✗	+/-	+/- +/-
Videoder-matoscope	Devices of different complexity and magnifications	✓	✓	✓	✓	↓	↑↑↑ ^a +/-
Confocal microscopy	Microscopy with high-resolution visualization of the superficial layers of the skin. Resolution similar to histopathology	✗	✓	✓	✓	↓↓	↑↑↑ ↑↑↑
Optical coherence tomography	Uses light sources (interferometry), obtaining non-invasive high-resolution imaging of the superficial structures of skin and mite	✗	✓	✓	✓	↓↓	↑↑↑ ↑↑↑
Optical microscopy	Visualization under optical microscope of samples obtained with adhesive tape (adhesive tape test), scraping (Müller test), or biopsy	✗	✓	✗	✗	+/- ^a	+/- ^a ↑ ^a
Serological and molecular tests	Detection of mite targets enabling simple and rapid point-of-care diagnosis					Currently not available	

✓, yes; ✗, no; ↑, high; ↑↑, very high; ↑↑↑, extremely high; ↓, low; ↓↓, too low; ±, medium.
Green color: positive data; red color: difficulty.

^a Differing for the different techniques available.

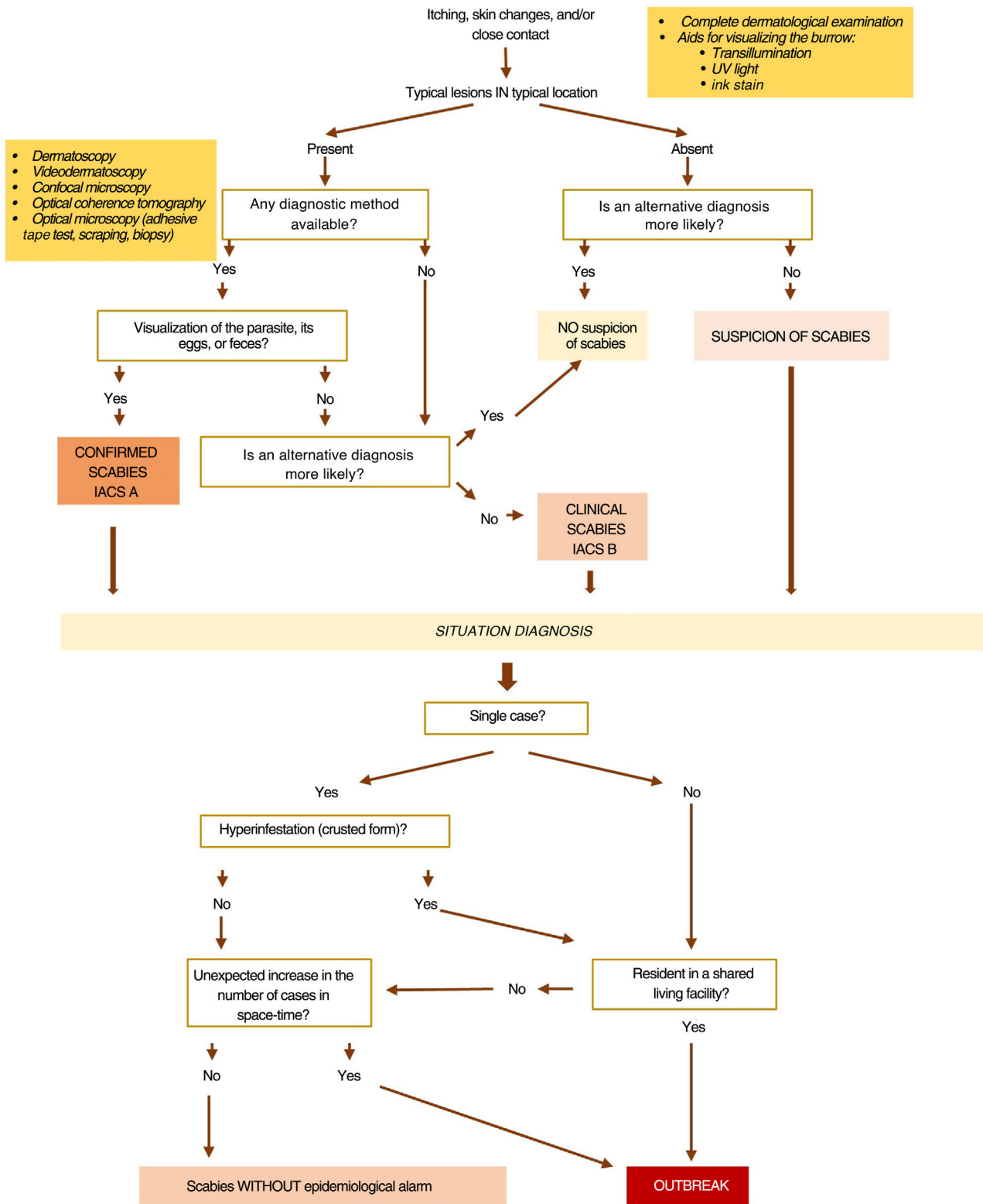


Figure 2 Diagnostic algorithm of scabies.

has been achieved in most severe cases (level of evidence ⁴, grade of recommendation C).¹⁰ Additionally, in the presence of crusts or hyperkeratosis, keratolytics should be added to the scabicides.^{12,14,68}

In high-risk patients, such as immunocompromised individuals with classical scabies, combined treatment from the beginning may be considered.^{67,69} (level of evidence 5, grade of recommendation D).

Table 4 Mode of application of topical scabicides.

Application areas	Observations
<i>Head and face:</i> avoiding periocular and perioral area	Only in children younger than 1 year and crusted forms
<i>Body:</i> cover from mandibular arch and retroauricular folds to tips of toes	In all cases
<i>Areas of special attention:</i> careful application in skin folds and periungual areas and less accessible areas	In all cases
Application method	
1. Trim nails before application.	
2. Remove crusts, as much as possible, before application, in crusted or hyperkeratotic forms.	
3. Apply to an unclothed person.	
4. Assistance from another person to make sure application covers the entire recommended skin area.	
5. When the helper does not need treatment because he/she is not a close contact and does not suffer from scabies, he/she should wear gloves and contact protection measures.	
6. Reapply the product on areas that were washed during the recommended drug contact time such as hands, diaper area...	
7. Rinse off the product in the shower after the recommended contact time has gone by.	
8. All affected individuals and contacts in the household must undergo scabicide treatment (it does not need to be the same drug) within a 24-hour window to prevent reinfections.	

Drug resistance

In recent years, therapeutic failures have been described with all scabicides.^{3,70–72} Contributing factors may include drug resistance generation,⁷³ inadequate cutaneous bioavailability of the drug,⁷⁴ incorrect treatment compliance, and reinfection due to insufficient contact control.^{7,75} Poor control of fomites does not seem to be a plausible cause, given the low frequency of this source of contagion and the demonstrated efficacy in community campaigns that do not involve environmental mite elimination.^{76,77}

In vitro tolerance of the mite to permethrin and ivermectin has increased since they started being used for scabies.^{71,78,79} In addition, mutations related to permethrin resistance have been identified. However, the few available clinical studies still report high efficacy rates.^{80,81}

It is difficult to prove the cause of failures for several reasons: there is no experimental culture medium for the mite, and in vitro studies lack standardization; confirmation of correct treatment compliance is not always possible, and, eventually, the higher the prevalence the more difficult it becomes to access to all individuals who need treatment as direct contacts.

In case of therapeutic failure, due to absence of cure criteria 4 weeks into treatment, the complete regimen should be repeated. After confirming compliance with all necessary measures—personal, contacts, and environmental—the health care worker must decide whether to repeat the previous regimen, extend the skin contact time with the topical scabicide, increase the number of cycles of oral or topical treatment, prescribe the association of permethrin and ivermectin, or recommend an alternative treatment, among those currently considered second-line therapies. This decision should take into account the preferences and possible contraindications of the affected individual and his/her contacts (figure 3).

Post-mite elimination treatment (persistent symptoms)

After mite elimination, skin lesions, itching, and bacterial complications may persist for 2 to 4 weeks.

Itching: oral antihistamines can alleviate it (level of evidence 5, grade of recommendation D). The topical application of medium-potency corticosteroids and tacrolimus has proven effective in reducing itching (level of evidence 1b, grade of recommendation A).⁸³ Before starting the application, the absence of bacterial superinfection and parasitological cure should be confirmed to avoid masking infection and persistent scabies.

Skin lesions: after parasitological cure, topical corticosteroids can help resolve the inflammatory process (level of evidence 5, grade of recommendation D).

Bacterial infections: they should be treated with topical or oral antibiotics based on their severity and, eventually, they should be antibiogram-guided (level of evidence 2c, grade of recommendation B).⁸⁴

Management of direct contacts

In cases of classical scabies, all close primary contacts of the affected individual from 1 to 2 months before the onset of symptoms, should be treated, regardless of the presence of symptoms, including sexual partners and individuals who share a bed or have close household or institutional contact (schools, shelters, nursing homes, etc.).

For secondary contacts—those who have had close contact with a primary contact of a case—follow-up is advised.

The sexual partners in the 2 months prior to symptom onset should be examined and treated (level of evidence 5, grade of recommendation D). Close contact should be avoided until the affected individual and his/her partners have been treated (level of evidence 5, grade of recommendation D).^{10,11}

In cases of hyperinfestation, all primary contacts should be treated, even if they have not been in intimate or prolonged contact. Secondary contacts should also be monitored, and treatment should be considered.^{12,62,81}

Table 5 Scabicides available in Spain.

Active ingredient	Commercial brands	Mechanism of action	Ovicidal activity	Therapeutic formulation	Dosage	Efficacy and recommendation	Adverse events	Pediatric use	Use in pregnancy and lactation	Price ^{†, ††}
Permethrin	Perme-cure cream Sarcop cream	Pyrethroid Neurotoxic Blocks Na ⁺ channels	Doubtful	5% cream	Topical application Repeat within 7 to 14 days	First choice in adults (E1a, RA) Equal efficacy to ivermectin (E1a, RA) ^{10,85,86}	High safety profile Dermatitis	First choice > 2 months (E1a, RA) ¹⁰ (safety not established for < 2 months)	First choice (2b, RB) ⁸⁷ FDA pregnancy category B. Non-teratogenic in animal models	70 g: €17.02
Oral ivermectin	Ivergalen Ivercare Ivermectin Teva	Macrocyclic lactone Neurotoxic Blocks Cl ⁻ channels	No	Oral route 200 µg/kg ^a	Single dose Repeat within 7 to 14 days Ingest with fatty food ^b	First choice in adults (E1a, RA) ^{10,67} Equal efficacy to permethrin (E1a, RA) More suitable for population-wide treatments ^{76,77}	Very rare: Neurotoxicity, hepatotoxicity, nephrotoxicity, toxic epidermal necrolysis, and Steven-Johnson syndrome	First choice for > 15kg Ill-advised in < 15 kg (safety not established)	Ill-advised in pregnancy (safety not established) FDA Pregnancy category C Teratogenic in rats	3 mg × 8 tabs: €34.96

Table 5 (Continued)

Active ingredient	Commercial brands	Mechanism of action	Ovicidal activity	Therapeutic formulation	Dosage	Efficacy and recommendation	Adverse events	Pediatric use	Use in pregnancy and lactation	Price ^{†, ††}
Topical ivermectin	Available in magistral formulation	Macrocyclic lactone Neurotoxic Blocks Cl-channels	No	1% lotion	Topical application Repeat within 7 to 14 days	Similar efficacy to topical permethrin (E1b, RA) ⁶⁷	High safety profile ⁶⁷ Dermatitis	Ill-advised (safety not established)	Ill-advised (safety not established)	100 mL: €50 (approx.)
Sulfur	Available in magistral formulation	Keratolytic and scabicial action	Yes	Vaseline, emulsion, or 10% and 5% lotion (in patients younger than 1 year)	Topical application for 3 consecutive days Repeat cycle after 7 days	Contradictory efficacy data ^{86,88,89}	Dermatitis Asteatotic eczema. Common mild adverse events No serious adverse events described	First choice in infants < 2 months (E5, RD)	Recommended (E5, RD) ¹²	200 g/mL: €50 (approx.)
Benzyl benzoate	Available in magistral formulation	Unknown	Unknown	Emulsion or lotion for 1 to 12 years: 5% to 10% >12 years: 10% to 25%	Topical application for 3 consecutive days Repeat cycle after 7 days	Comparable efficacy to oral ivermectin (E1b, RA) ^{87,90}	Intense dermatitis, conjunctivitis ^{14,86}	Ill-advised in patients < 1 year ⁹¹	Recommended in pregnancy (E2b, RB) ⁹² Ill-advised in lactation (E4, RC) ^{10,93}	200 mL: €70 (approx.)

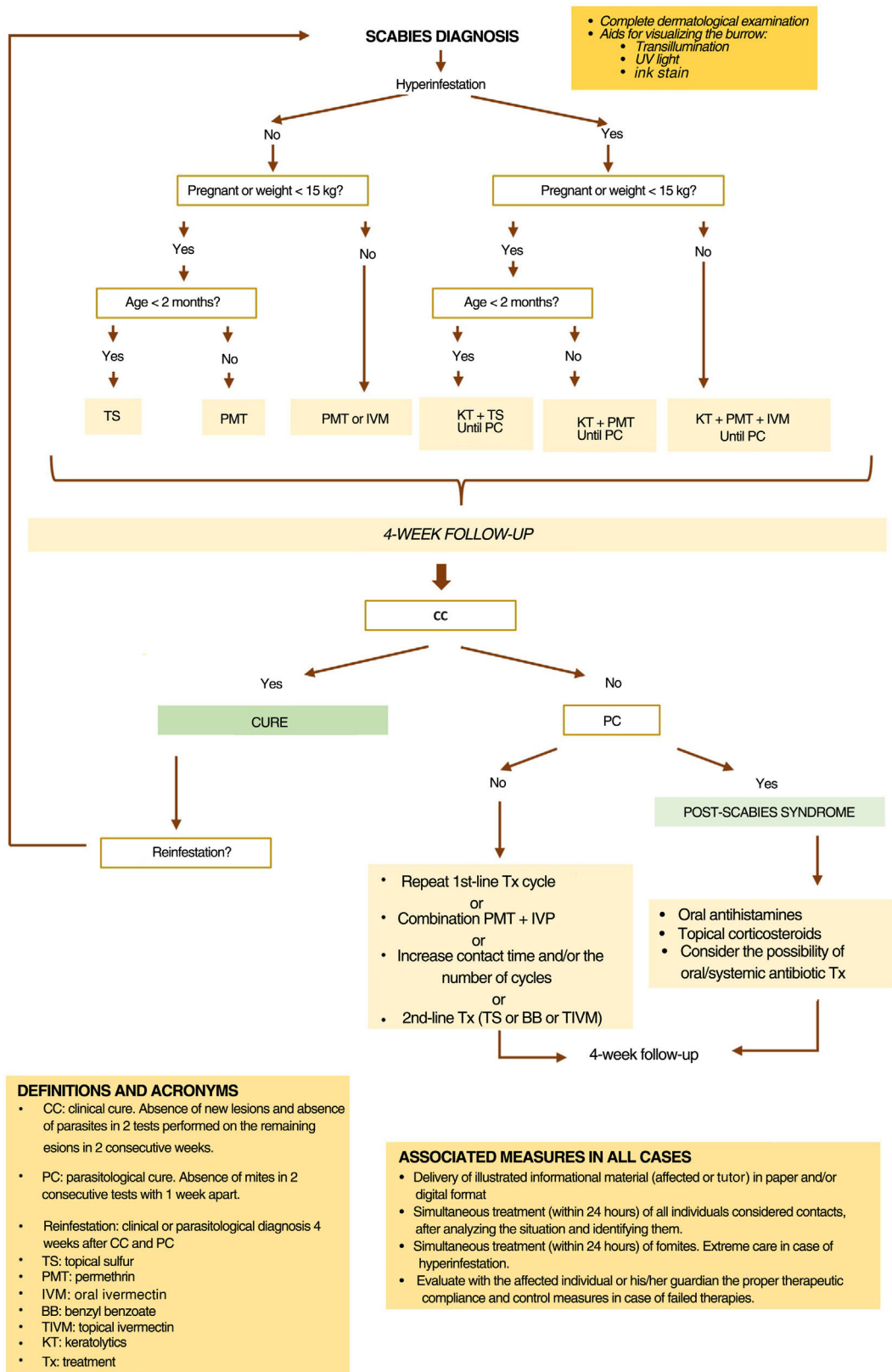
E, level of evidence; FDA, Food and Drug Administration; R, degree of recommendation.

^a Dosage table by weight in Annex 3.

^b Recommended to enhance absorption, unlike fasting intake, indicated for other diseases in the technical sheet.¹⁰

[†] Price for the recommended regimen for an adult of average build. In topical preparations, calculated based on 35 g per application. Figures from the Spanish Ministry of Health for drugs included in the pharmaceutical provision of the National Health System consulted in September 2023.²⁹

^{††} Approximate price calculated based on current documents on billing of magistral formulation from various Health Departments across various Spanish Autonomous Communities.
Green color: first choice; red color: ill-advised,



4-week follow-up

DEFINITIONS AND ACRONYMS

- CC: clinical cure. Absence of new lesions and absence of parasites in 2 tests performed on the remaining lesions in 2 consecutive weeks.
- PC: parasitological cure. Absence of mites in 2 consecutive tests with 1 week apart.
- Reinfestation: clinical or parasitological diagnosis 4 weeks after CC and PC
- TS: topical sulfur
- PMT: permethrin
- IVM: oral ivermectin
- BB: benzyl benzoate
- TIVM: topical ivermectin
- KT: keratolytics
- Tx: treatment

ASSOCIATED MEASURES IN ALL CASES

- Delivery of illustrated informational material (affected or tutor) in paper and/or digital format
- Simultaneous treatment (within 24 hours) of all individuals considered contacts, after analyzing the situation and identifying them.
- Simultaneous treatment (within 24 hours) of fomites. Extreme care in case of hyperinfestation.
- Evaluate with the affected individual or his/her guardian the proper therapeutic compliance and control measures in case of failed therapies.

Figure 3 Therapeutic algorithm of scabies.

Table 6 Pubic lice infestation. Diagnosis and control.

Clinical signs			
Diagnosis	Visualization of the parasite or its eggs (naked eye, magnifying glass, or dermatoscope)		
Symptom	Itching		
Dermatological signs	<ul style="list-style-type: none"> - Bluish spots (violaceous macules < 1 cm on the skin where the parasite fed from) - Erythematous papules - Ochre spots (parasite feces on the host's skin) 		
General guidelines			
<p>Give oral and written information on the disease, its treatment, and control measures (E4, RC). Investigate concomitant STIs (E2b, RB).^{94,95} Apply treatment to the affected hairy areas (pubis and other affected areas such as perianal, axillae, trunk, eyebrows, eyelashes, beard, and mustache). Use creams or lotions, preferably over shampoos. Repeat the treatment within 3 to 7 days (E2b, RB). Remove nits with hands or a fine-toothed comb. Decontaminate fomites (personal clothing, bedding, and towels) by washing in a hot water cycle followed by drying or sealing them in a bag for 72 hours. Review sexual partners from the previous 3 months and treat affected individuals and their sexual partners from the previous month (E4, RC). Avoid sharing clothing and bodily contact until the case and sexual partners have completed treatment. Perform clinical surveillance 1 week after treatment (E2b, RB).</p>			
Criterion for cure: absence of lice and/or viable eggs 1 week after completing treatment.			
Therapeutic measures			
Recommendation	Drug	Indication	Application method
First-line therapy	1% permethrin (scalp cream or lotion)	Indicated in all ages and in pregnant or breastfeeding women	Apply to damp area, rinse off in 10 minutes (E1b, RA) ⁹
Second-line therapy	Oral ivermectin 250 µg/kg	Not recommended in pregnancy and children under 15 kg	Ingest with fatty food. Repeat dose within 7 to 14 days (E5, RD) ⁹⁶
	Topical 1% ivermectin	Safety not studied in pregnant women or children	Topical application. No established regimen for this indication (E5, RD)
Not included in guidelines	25% benzyl benzoate lotion	Not recommended for children younger than 1 year-old	Topical application. No established regimen for this indication (E5, RD)
	Dimethicone	No studies on pubic lice. Effective in head lice and safe in pregnancy, breastfeeding, and children	Topical application. Regimen not described for this indication
	Shaving	It is not a pesticide; it works by suffocation. No risk of resistance. Efficient, cost-effective, and safe. Eliminates infestation by removing the lice's natural habitat. May be poorly accepted depending on location (eyelashes...), extent of infestation, and individual preferences	Removal of the portion of hair that emerges from the skin (it is not necessary to remove it from the root)

Table 6 (Continued)

Recommendation	Drug	Therapeutic measures	
		Indication	Application method
Treatment on eyelashes	Smooth petroleum jelly or ophthalmic ointment	No contraindications It acts by suffocation	Apply thick layer, in occlusive dressing (covered), twice daily for 10 days (E5, RD)
	Oral ivermectin	Not recommended in pregnancy and children under 15 kg	Same regimen and evidence
	1% permethrin	Safe	Apply with eyes closed, remove in 10 minutes (E5, RD)
	Mechanical removal of lice	Safe It may be necessary to trim some eyelashes	Removal with tweezers, visualization with slit lamp. More difficult for nits than hatched lice. Often requires subsequent treatment with petroleum jelly or ointment
Not currently available in Spain for human use	0.5% malathion lotion		
	Pyrethrins with topical piperonyl butoxide (E1b, RA) ¹¹		
	0.2% phenothrin lotion (E1b, RA)		
	0.5% to 1% carbaryl (E5, RD)		
	Lindane		

E, level of evidence; R, grade of recommendation.

Environmental mite elimination

For classical scabies, there is no evidence that fomite treatment is helpful.³⁸ However, based on studies that found mites in the environment,³⁸ most clinical practice guidelines recommend eliminating mites on fomites that have come into contact with the affected individual's skin (level of evidence 5, grade of recommendation D).¹⁰

Mites can be completely eliminated by exposing clothes to 50°C temperatures for 10 minutes (in washing machine, tumble dryer or water), keeping them in sealed bags for 3 (moderate ambient temperature and humidity) to 8 days (high ambient humidity and low temperatures), or keeping them in the freezer at -10°C for 5 hours.⁹⁷ Surfaces should be cleaned with soap, water, and a vacuum cleaner.¹⁷

To avoid reinfections, these measures should be implemented the same day as the treatment for affected individuals and contacts.

For crusted scabies, where patients harbor a very high number of mites, all guidelines recommend rigorous environmental decontamination.

Outbreak control (Figure 2 and supplementary data. Annexes 4 and 5)

In the event of an institutional outbreak of scabies, the following measures should be implemented:

- Notify the administration and those responsible for prevention and occupational health at the center, as well as

residents, their families, and the centers the residents may have recently been transferred to.

- Provide basic training on the disease to workers.
- Daily evaluation of symptoms and signs in residents, workers, and new admissions while the outbreak persists.
- Treatment of those meeting scabies diagnostic criteria.
- Offer preventive treatment to workers, visiting family members, and asymptomatic residents living in the affected area of the institution.
- Temporary work disability for affected workers for as long as they exhibit signs of active infestation and, at least, 24 hours after the first dose of treatment.
- Contact isolation measures for affected residents.
- Implementation of measures for environmental mite elimination.
- Detailed notification of the situation to regional health authorities.
- These same measures should be implemented to deal with cases of crusted scabies, and all close primary contacts should be treated as well. Secondary contacts and individuals in contact with the affected individual's fomites, such as laundry personnel, should be evaluated and offered curative or preventive treatment.^{15,17,19,62}

Pubic lice

Pubic lice, also known as "crab louse", are caused by the ectoparasite *Phthirus pubis*. They are visible to the naked eye

and smaller than head lice. They infest pubic hair and, to a lesser extent, body hair, eyebrows, eyelashes and armpits.

Transmission and epidemiology

Pubic lice are primarily transmitted through sexual contact, but depending on the site of infestation, they can also spread through non-sexual contact among close household contacts. Transmission via fomites is much less likely, as the lice cannot survive for 24 hours without feeding from the host's blood.

The eggs are deposited, attached to the hair, close to the skin. After hatching, typically a week after being laid, the empty eggshell may remain attached to the hair. Nymphs reach maturity and become capable of contagion within 2 weeks.⁹⁴

Currently, there are no current prevalence studies in the adult population. Various studies, albeit not recent, evaluate cases seen in STI clinics with prevalence rates close to 2%.⁹⁵

The incidence of pubic lice is increasing in disadvantaged communities with overcrowded conditions and decreasing in populations where pubic hair removal is a common practice. Affected individuals are more likely than the general population to have other concurrent STIs.^{8,9}

Clinical presentation, treatment, and control

Table 6 summarizes the clinical characteristics and recommended treatment and control measures.

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Conflicts of interest

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at [doi:10.1016/j.ad.2024.03.016](https://doi.org/10.1016/j.ad.2024.03.016).

References

- Redondo-Bravo L, Fernandez-Martinez B, Gómez-Barroso D, Gherasim A, García-Gómez M, Benito A, et al. Scabies in Spain? A comprehensive epidemiological picture. *PLoS One*. 2021;16:e0258780, <http://dx.doi.org/10.1371/journal.pone.0258780>.
- Louka C, Logothetis E, Engelman D, Samiotaki-Logotheti E, Pournaras S, Stienstra Y. Scabies epidemiology in health care centers for refugees and asylum seekers in Greece. *PLoS Negl Trop Dis*. 2022;16:e0010153, <http://dx.doi.org/10.1371/journal.pntd.0010153>.
- Mazzatenta C, Piccolo V, Argenziano G, Bassi A. Is Scabies becoming less sensitive to permethrin therapy? *J Eur Acad Dermatol Venereol*. 2021;35, <http://dx.doi.org/10.1111/jdv.17339>.
- Martínez-Pallás I, Aldea-Manrique B, Ramírez-Lluch M, Manuel Vinuesa-Hernando J, Ara-Martín M. Scabies outbreak during home confinement due to the SARS-CoV-2 pandemic. *J Eur Acad Dermatol Venereol*. 2020;34, <http://dx.doi.org/10.1111/jdv.16879>.
- Cerro PA, Navarro-Bielsa A, Palma AM. FR - Epidemia de sarna en el contexto de la pandemia de COVID-19. *Actas Dermosifiliogr*. 2022;113:516–8, <http://dx.doi.org/10.1016/j.ad.2020.11.028>.
- Aguado Vázquez Á, Gegúndez Hernández H, Melgosa Ramos FJ, Díaz Corpas T. Prevalencia y características clínicas de pacientes diagnosticados de escabiosis durante la pandemia producida por el coronavirus de tipo 2 causante del síndrome respiratorio agudo (SARS-CoV-2) en un hospital de tercer nivel. Un estudio descriptivo. *Actas Dermosifiliogr*. 2023;114:171–2, <http://dx.doi.org/10.1016/j.ad.2022.05.017>.
- Galván Casas C, Ruiz Villaverde C, Prados Carmona Á, Fernández Camporro A, Angulo Menéndez AG, Álvarez-Buylla Puente MC, et al. Características clínicas y epidemiológicas de pacientes diagnosticados de escabiosis en España: oportunidades de mejora. Estudio transversal multicéntrico CLINI-AEDV. *Actas Dermosifiliogr*, <https://doi.org/10.1016/j.ad.2023.08.006>.
- Pierzchalski JL, Bretl DA, Matson SC. Phthirus pubis as a Predictor for Chlamydia Infections in Adolescents. *Sex Transm Dis*. 2002;29:331–4, <http://dx.doi.org/10.1097/00007435-200206000-00004>.
- Salavastru CM, Chosidow O, Janier M, Tiplica GS. European guideline for the management of pediculosis pubis. *J Eur Acad Dermatol Venereol*. 2017;31:1425–8, <http://dx.doi.org/10.1111/jdv.14420>.
- Salavastru CM, Chosidow O, Boffa MJ, Janier M, Tiplica GS. European guideline for the management of scabies. *J Eur Acad Dermatol Venereol*. 2017;31:1248–53, <http://dx.doi.org/10.1111/jdv.14351>.
- Tiplica GS, Radcliffe K, Evans C, Gomberg M, Nandwani R, Rafila A, et al. 2015 European guidelines for the management of partners of persons with sexually transmitted infections. *J Eur Acad Dermatol Venereol*. 2015;29:1251–7, <http://dx.doi.org/10.1111/jdv.13181>.
- Executive Committee of Guideline for the Diagnosis and Treatment of Scabies. Guideline for the diagnosis and treatment of scabies in Japan (third edition); Executive Committee of Guideline for the Diagnosis and Treatment of Scabies. *J Dermatol*. 2017;44:991–1014, <http://dx.doi.org/10.1111/1346-8138.13896>.
- Sunderkötter C, Feldmeier H, Fölster-Holst R, Gomberg M, Nandwani R, Rafila A, et al. S1 guidelines on the diagnosis and treatment of scabies - Short version. *J Dtsch Dermatol Ges*. 2016;14:1155–67, <http://dx.doi.org/10.1111/ddg.13130>.
- Sashidharan PM, Basavaraj S, Bates CM. 2016 UK National Guideline on the Management of Scabies. Published online March 1. 2016 [accessed 22 Aug 2023]. Available from:

- <https://www.bashguidelines.org/media/1137/scabies-2016.pdf>
15. Prevention CC for DC and. CDC - Scabies - Resources for Health Professionals - Institutional Settings. Published April 19, 2019. Accessed August 27, 2023. [accessed 29 Aug 2023]. Available from: https://www.cdc.gov/parasites/scabies/health_professionals/institutions.html
 16. Workowski KA, Bachmann LH, Chan PA, Johnston CM, Muzny CA, Park I, et al. Sexually Transmitted Infections Treatment Guidelines. Published online July 23, 2021. Available from: <https://www.cdc.gov/std/treatment-guidelines/STI-Guidelines-2021.pdf>
 17. Barrabeig I, Gassiot M, Gómez de Carvallo M, Maresma MM, Molinero E, Parrón I, et al. Subdirecció General de Vigilància i Resposta a Emergències, de Salut Pública. Protocol de prevenció i control de l'escabiosi (sarna). 2023 [accessed 22 Aug 2023]. Available from: https://scientiasalut.gencat.cat/bitstream/handle/11351/4203.2/protocol_prevencio_control_escabiosi_sarna_2023pdf?sequence=8&isAllowed=5
 18. Casado Casuso S, Borrego Izquierdo Y, Prieto Sánchez R, Oro Fernández M. Actualización del tratamiento de la sarna (Cantabria). Published online. 2021 [accessed 22 Aug 2023]. Available from: <https://www.scsalud.es/documents/2162705/2163020/2021+01+Ivermectina+para+la+sarna.pdf/b0d0d4d7-8fbc-587f-1757-69b78ad5397e?t=161166200094>
 19. Consejería de Salud - Dirección General de Salud Pública y Adicciones. Servicio de Epidemiología. Gobierno de Murcia. Protocolo de actuación frente a escabiosis. Published online March 2022. [accessed 22 Aug 2023]. Available from: https://www.murciasalud.es/recursos/ficheros/499065-Protocolo_escabiosis_DEF_2022.pdf#:~:text=El%20paciente%20diagnosticado%20de%20sarna%20deber%C3%A1%20realizar%20el%20siguiente%20tratamiento%3A&text=Se%20recomienda%20duchar%206%20minutos,u%C3%B1as%20de%20manos%20y%20pies.&text=producto%20mediante%20una%20ducha%20de,deber%C3%A1%20estar%20desinfectada%20y%20limpia
 20. Departamento de Salud del Gobierno Vasco. Tratamiento de la sarna. Published online 2022. [accessed 22 Aug 2023]. Available from: https://www.euskadi.eus/contenidos/informacion/cevime_infac_2022/es_def_adjuntos/INFAC_Vol_30_3_SARNA.pdf
 21. Xunta de Galicia. Protocolo de actuación ante a sarna humana en institucións con cuidadores. Published online November 2017. [accessed 22 Aug 2023]. Available from: https://www.sergas.es/Saude-publica/Documents/1312/Protocolo_sarna_2017.pdf
 22. Servicio de Vigilancia Epidemiológica de Andalucía. Protocolo de intervención ante alerta por infestación por ectoparasitos Versión 2. Documento no publicado.
 23. Drake Monfort M, Abraira Meriel C, Naharro Fernandez C, Lopez Sundh AE, González López MA. Protocolo diagnóstico terapéutico de escabiosis Servicio de Dermatología Hospital Universitario Marqués de Valdecilla. Diciembre 2021. Documento no publicado.
 24. Departamento de Salud Gobierno Vasco. Unidad de vigilancia epidemiológica. Protocolo de actuación ante casos de escabiosis. 8 abril 2022. Documento no publicado.
 25. Red de Vigilancia Epidemiológica de la Comunidad de Madrid. Protocolo de actuación frente a la escabiosis. https://www.comunidad.madrid/sites/default/files/doc/sanidad/epid/protocolo_escabiosis_octubre2022.pdf
 26. Gerencia de Atención Primaria. Tratamiento y medidas higiénicas de la escabiosis. [accessed 22 Aug 2023]. Available from: <https://www.ibsalut.es/apmallorca/es/profesionales/publicaciones/consejos-de-salud/seguridad-y-prevencion-del-riesgo/2209-sarna-tratamiento-y-medidas-higienicas>
 27. Dirección General de Salud Pública. Protocolo escabiosis o sarna. Published online July 2023. [accessed 22 Aug 2023]. Available from: https://www.riojasalud.es/files/content/salud-publica-consumo/epidemiologia/alertas/sarna/Protocolo_La-Rioja_Sarna.pdf
 28. OCEBM Levels of Evidence Working Group. The Oxford Levels of Evidence. Published online March 2009. [accessed 28 Aug 2023]. Available from: <https://www.cebm.ox.ac.uk/resources/levels-of-evidence/oxford-centre-for-evidence-based-medicine-levels-of-evidence-march-2009>
 29. Ministerio de Sanidad - Profesionales de la Salud - Nomenclator [accessed 27 Aug 2023]. Available from: <https://www.sanidad.gob.es/profesionales/nomenclator.do>
 30. Arlian LG, Morgan MS. A review of Sarcoptes scabiei: Past, present and future. Parasit Vectors. 2017;10:297, <http://dx.doi.org/10.1186/s13071-017-2234-1>.
 31. Puza CJ, Suresh V. Scabies and Pruritus. A Historical Review. JAMA Dermatol. 2018;154:536, <http://dx.doi.org/10.1001/jamadermatol.2018.0147>.
 32. Walton SF, Holt DC, Currie BJ, Kemp DJ. Scabies: New Future for a Neglected Disease In: Advances in Parasitology, 57. Elsevier; 2004. p. 309–76, [http://dx.doi.org/10.1016/S0065-308X\(04\)57005-7](http://dx.doi.org/10.1016/S0065-308X(04)57005-7).
 33. Liu JM, Wang HW, Chang FW, Kemp DJ. The effects of climate factors on scabies. A 14-year population-based study in Taiwan. Parasite. 2016;23:54, <http://dx.doi.org/10.1051/parasite/2016065>.
 34. Golant AK, Levitt JO. Scabies: A Review of Diagnosis and Management Based on Mite Biology. Pediatr Rev. 2012;33:e1–12, <http://dx.doi.org/10.1542/pir.33.1.e1>.
 35. Arlian LG, Runyan RA, Achar S, Estes SA. Survival and infestivity of Sarcoptes scabiei var. canis and var. hominis. J Am Acad Dermatol. 1984;11:210–5, [http://dx.doi.org/10.1016/S0190-9622\(84\)70151-4](http://dx.doi.org/10.1016/S0190-9622(84)70151-4).
 36. Fuller LC. Epidemiology of scabies. Curr Opin Infect Dis. 2013;26:123–6 <https://doi.org/10.1097/qco.0b013e32835eb851>
 37. Van Der Linden N, van Gool K, Gardner K, Dickinson H, Agostino J, Regan DG, et al. A systematic review of scabies transmission models and data to evaluate the cost-effectiveness of scabies interventions. PLoS Negl Trop Dis. 2019;13:e0007182 <https://doi.org/10.1371/journal.pntd.0007182>
 38. Mellanby K. Transmission of Scabies. Br Med J. 1941;2:405–6 <https://doi.org/10.1136/Fbmj.2.4211.405>
 39. Mellanby K. The development of symptoms, parasitic infection and immunity in human scabies. Parasitology. 1944;35:197–206, http://dx.doi.org/10.1017/S_0031182000021612.
 40. Belvisi V, Orsi GB, del Borgo C, Fabiatti P, Ianari A, Albertoni F, et al. Large Nosocomial Outbreak Associated with a Norwegian Scabies Index Case Undergoing TNF- α Inhibitor Treatment: Management and Control. Infect Control Hosp Epidemiol. 2015;36:1358–60 <https://doi.org/10.1017/ice.2015.188>
 41. Engelman D, Marks M, Steer AC, Beshah A, Biswas G, Chosidow O, et al. A framework for scabies control. PLoS Negl Trop Dis. 2021;15:e0009661, <http://dx.doi.org/10.1371/journal.pntd.0009661>.
 42. Andriantsoanirina V, Izri A, Botterel F, Foulet F, Chosidow O, Durand R. Molecular survey of knock-down resistance to pyrethroids in human scabies mites. Clin Microbiol Infect. 2014;20:O139–41, <http://dx.doi.org/10.1111/1469-0691.12334>.
 43. Moroni B, Rossi L, Bernigaud C, Guillot J. Zoonotic Episodes of Scabies: A Global Overview. Pathogens. 2022;11:213, <http://dx.doi.org/10.3390/pathogens11020213>.
 44. Aydingöz İE, Mansur AT. Canine Scabies in Humans: A Case Report and Review of the Literature. Dermatology. 2011;223:104–6, <http://dx.doi.org/10.1159/000327378>.

45. Chosidow O. Scabies and pediculosis. *Lancet*. 2000;355:819–26, [http://dx.doi.org/10.1016/S0140-6736\(99\)09458-1](http://dx.doi.org/10.1016/S0140-6736(99)09458-1).
46. Engelman D, Cantey PT, Marks M, Solomon AW, Chang AY, Chosidow O, et al. The public health control of scabies: Priorities for research and action. *Lancet*. 2019;394:81–92, [http://dx.doi.org/10.1016/S0140-6736\(19\)31136-5](http://dx.doi.org/10.1016/S0140-6736(19)31136-5).
47. WHO. Scabies. Published online May 31, 2023. [accessed 12 Oct 2023]. Available from: <https://www.who.int/news-room/fact-sheets/detail/scabies>
48. World Health Organization. Ending the Neglect to Attain the Sustainable Development Goals: A Sustainability Framework for Action against Neglected Tropical Diseases 2021–2030. World Health Organization; 2021. [accessed 29 Aug 2023]. Available from: <https://apps.who.int/iris/handle/10665/338886>
49. Lydeamore MJ, Campbell PT, Regan DG, Tong SYC, Andrews RM, Steer AC, et al. A biological model of scabies infection dynamics and treatment informs mass drug administration strategies to increase the likelihood of elimination. *Math Biosci*. 2019;309:163–73, <http://dx.doi.org/10.1016/j.mbs.2018.08.007>.
50. Kinyanjui T, Middleton J, Güttel S, Cassell J, Ross J, House T. Scabies in residential care homes: Modelling, inference and interventions for well-connected population sub-units. *PLoS Comput Biol*. 2018;14:e1006046, <http://dx.doi.org/10.1371/journal.pcbi.1006046>.
51. Thornton J. Scabies in Cox's Bazar. *Lancet*. 2023;402:600, [http://dx.doi.org/10.1016/S0140-6736\(23\)01731-2](http://dx.doi.org/10.1016/S0140-6736(23)01731-2).
52. Fernández Camporro Á, Navarro Fernández Í, Arcos González P. Escabiosis en España: tendencias del interés público y consumo de ectoparasiticidas. *Actas Dermosifiliogr*. 2023, <http://dx.doi.org/10.1016/j.ad.2022.11.018>. S0001731023005239.
53. Cox V, Fuller LC, Engelman D, Steer A, Hay RJ. Estimating the global burden of scabies: What else do we need? *Br J Dermatol*. 2021;184:237–42, <http://dx.doi.org/10.1111/bjd.19170>.
54. Martínez-García E, Grau-Pérez M, Buendía-Eisman A, García-Doval I. Prescriptions for scabies are rapidly increasing in Spain: An ecological study with national prescription data, 2008–2021. *J Eur Acad Dermatol Venereol*. 2023;37, <http://dx.doi.org/10.1111/jdv.18599>.
55. Thomas C, Coates SJ, Engelman D, Chosidow O, Chang AY. Ectoparasites: Scabies. *J Am Acad Dermatol*. 2020;82:533–48, <http://dx.doi.org/10.1016/j.jaad.2019.05.109>.
56. Pakanati K, Jagota D, Ladogana M. Norwegian scabies in HIV/AIDS. *Bayl Univ Med Cent Proc*. 2022;35:346–7, <http://dx.doi.org/10.1080/08998280.2022.2028705>.
57. Niode NJ, Adji A, Gazpers S, Kandou RT, Pandaleke H, Trisnowati DM, et al. Crusted Scabies, a Neglected Tropical Disease: Case Series and Literature Review. *Infect Dis Rep*. 2022;14:479–91, <http://dx.doi.org/10.3390/idr14030051>.
58. Mitchell E, Bell S, Thean LJ, Sahukhan A, Kama M, Koroivueti A, et al. Community perspectives on scabies, impetigo and mass drug administration in Fiji: A qualitative study. *PLoS Negl Trop Dis*. 2020;14:e0008825, <http://dx.doi.org/10.1371/journal.pntd.0008825>.
59. Engelman D, Yoshizumi J, Hay RJ, Osti M, Micali G, Norton S, et al. The 2020 International Alliance for the Control of Scabies Consensus Criteria for the Diagnosis of Scabies. *Br J Dermatol*. 2020;183:808–20, <http://dx.doi.org/10.1111/bjd.18943>.
60. Mundhra R, Rambhia K, Makhecha M, Gera R. Dermoscopy of crusted scabies: Revisiting the new and old signs. *Indian Dermatol Online J*. 2023;14:279, <http://dx.doi.org/10.4103/idoj.idoj.263.22>.
61. Micali G, Lacarrubba F, Verzi AE, Chosidow O, Schwartz RA. Scabies: Advances in Noninvasive Diagnosis. *PLoS Negl Trop Dis*. 2016;10:e0004691, <http://dx.doi.org/10.1371/journal.pntd.0004691>.
62. English LT, Terashita D, Kamali T, Cho K, Baron M, Pandes L, et al. Scabies prevention and control guidelines for health-care settings. 2019 [accessed 22 Aug 2023]. Available from: <http://www.publichealth.lacounty.gov/acd/Diseases/Scabies.htm>
63. Sunderkötter C, Wohlrab J, Hamm H. Scabies: Epidemiology, diagnosis, and treatment. *Dtsch Arztebl Int*. 2021;118:695–704, <http://dx.doi.org/10.3238/arztebl.m2021.0296>.
64. Mounsey KE, Bernigaud C, Chosidow O, McCarthy JS. Prospects for Moxidectin as a New Oral Treatment for Human Scabies. *PLoS Negl Trop Dis*. 2016;10:e0004389, <http://dx.doi.org/10.1371/journal.pntd.0004389>.
65. Cotreau MM, Warren S, Ryan JL, Fleckenstein L, Vanapalli SR, Brown KR, et al. The Antiparasitic Moxidectin: Safety Tolerability, and Pharmacokinetics in Humans. *J Clin Pharmacol*. 2003;43:1108–15, <http://dx.doi.org/10.1177/0091270003257456>.
66. Santos VSV, Pereira BB. Properties, toxicity and current applications of the biolarvicide spinosad. *J Toxicol Environ Health Part B*. 2020;23:13–26, <http://dx.doi.org/10.1080/10937404.2019.1689878>.
67. Thadanipon K, Anothaisintawee T, Rattanasiri S, Thakkintian A, Attia J. Efficacy and safety of antiscabietic agents: A systematic review and network meta-analysis of randomized controlled trials. *J Am Acad Dermatol*. 2019;80:1435–44, <http://dx.doi.org/10.1016/j.jaad.2019.01.004>.
68. Communicable Disease Section. Scabies control guidelines. Published online October 2015. [accessed 12 Oct 2023]. Available from: <https://www.health.vic.gov.au/infectious-diseases/scabies-control-guidelines>
69. Ichikawa M, Tanaka M, Naritomi Y, Furue M. Combined ivermectin and topical therapy significantly reduces treatment time in aged scabietic patients. *J Dermatol*. 2013;40:306–7, <http://dx.doi.org/10.1111/1346-8138.12070>.
70. Mounsey KE, Holt DC, McCarthy JS, Currie BJ, Walton SF. Longitudinal evidence of increasing in vitro tolerance of scabies mites to ivermectin in scabies-endemic communities. *Arch Dermatol*. 2009;145:840–1, <http://dx.doi.org/10.1001/archdermatol.2009.125>.
71. Currie BJ, Harumal P, McKinnon M, Walton SF. First documentation of in vivo and in vitro ivermectin resistance in *Sarcoptes scabiei*. *Clin Infect Dis Off Publ Infect Dis Soc Am*. 2004;39:e8–12, <http://dx.doi.org/10.1086/421776>.
72. Pasay C, Arlian L, Morgan M, et al. The Effect of Insecticide Synergists on the Response of Scabies Mites to Pyrethroid Acaricides. *PLoS Negl Trop Dis*. 2009;3:e354, <http://dx.doi.org/10.1371/journal.pntd.0000354>.
73. Riebenbauer K, Purkhauser K, Walochnik J, Urban N, Weber PB, Stamm T, et al. Detection of a knockdown mutation in the voltage-sensitive sodium channel associated with permethrin tolerance in *Sarcoptes scabiei* var. *hominis* mites. *J Eur Acad Dermatol Venereol*. 2023;37:2355–61, <http://dx.doi.org/10.1111/jdv.19288>.
74. Scholz L, Fritz C, Chuttke J, Eichner A, Wohlrab J. Permethrin Steal Effect by Unmasked Corneocytic Keratin in Topical Therapy of Scabies. *Skin Pharmacol Physiol*. 2023;36:107–16, <http://dx.doi.org/10.1159/000529401>.
75. Nemecek R, Stockbauer A, Lexa M, Poepl W, Moeseder G. Application errors associated with topical treatment of scabies: An observational study. *J Dtsch Dermatol Ges*. 2020;18:554–9, <http://dx.doi.org/10.1111/ddg.14122>.
76. Romani L, Marks M, Sokana O, Nasi T, Kamoriki B, Cordell B, et al. Efficacy of mass drug administration with ivermectin for control of scabies and impetigo, with coadministration of azithromycin: A single-arm community intervention trial. *Lancet Infect Dis*. 2019;19:510–8, [http://dx.doi.org/10.1016/S1473-3099\(18\)30790-4](http://dx.doi.org/10.1016/S1473-3099(18)30790-4).

77. Galván-Casas C, Mitjá O, Esteban S, Kafulafula J, Phiri T, Navarro-Fernández Í, et al. A facility and community-based assessment of scabies in rural Malawi. *PLoS Negl Trop Dis*. 2021;15:e0009386, <http://dx.doi.org/10.1371/journal.pntd.0009386>.
78. Fraser J. Permethrin: A Top End viewpoint and experience. *Med J Aust*. 1994;160:806-806, <http://dx.doi.org/10.5694/j.1326-5377.1994.tb125968.x>.
79. Walton SF, Myerscough MR, Currie BJ. Studies in vitro on the relative efficacy of current acaricides for *Sarcoptes scabiei* var. *hominis*. *Trans R Soc Trop Med Hyg*. 2000;94:92-6, [http://dx.doi.org/10.1016/S0035-9203\(00\)90454-1](http://dx.doi.org/10.1016/S0035-9203(00)90454-1).
80. Yürekli A. Is there a really resistance to scabies treatment with permethrin? In vitro killing activity of permethrin on *Sarcoptes scabiei* from patients with resistant scabies. *Dermatol Ther*. 2022;35:e15260, <http://dx.doi.org/10.1111/dth.15260>.
81. Sunderkötter C, Aebischer A, Neufeld M, Löser C, Kreuter A, Bialek R, et al. Increase of scabies in Germany and development of resistant mites? Evidence and consequences. *J Dtsch Dermatol Ges*. 2019;17:15-23, <http://dx.doi.org/10.1111/ddg.13706>.
82. Yürekli A, Can İ, Oğuz M. Using ultraviolet light in diagnosing scabies: Scabies' Sign via Wood's Lamp. *J Am Acad Dermatol*. 2023;89:e195-6, <http://dx.doi.org/10.1016/j.jaad.2023.07.006>.
83. Yadav P, Mohan S, Sonthalia S, Ramesh V, Kashyap V. A comparative study of topical tacrolimus and topical triamcinolone acetonide in nodular scabies. *Dermatol Ther*. 2020;33:e13954, <http://dx.doi.org/10.1111/dth.13954>.
84. Chung SD, Wang KH, Huang CC, Lin HC. Scabies increased the risk of chronic kidney disease: A 5-year follow-up study. *J Eur Acad Dermatol Venereol*. 2014;28:286-92, <http://dx.doi.org/10.1111/jdv.12099>.
85. Rosumeck S, Nast A, Dressler C. Ivermectin and permethrin for treating scabies Cochrane Infectious Diseases Group. *Cochrane Database Syst Rev*. 2018;2018, <http://dx.doi.org/10.1002/14651858.CD012994>.
86. Strong M, Johnstone P. Interventions for treating scabies. *Cochrane Database Syst Rev*. 2007;2007, <http://dx.doi.org/10.1002/14651858.CD000320.pub2>. CD000320.
87. Porto I. Antiparasitic Drugs and Lactation: Focus on Anthelmintics Scabicides, and Pediculicides. *J Hum Lact*. 2003;19:421-5, <http://dx.doi.org/10.1177/0890334403258133>.
88. Chhaiya S, Dave J, Shah H, Patel V, Mehta D. Comparative efficacy and safety of topical permethrin, topical ivermectin, and oral ivermectin in patients of uncomplicated scabies. *Indian J Dermatol Venereol Leprol*. 2012;78:605, <http://dx.doi.org/10.4103/0378-6323.100571>.
89. Fathy FM, El-Kasah F, El-Ahwal AM. Clinical and parasitological study on scabies in Sirte, Libya. *J Egypt Soc Parasitol*. 2010;40:707-31.
90. Ertugrul G, Aktas H. Comparison of sulfur ointment and permethrin treatments in scabies. *Dermatol Ther*. 2022;35, <http://dx.doi.org/10.1111/dth.15897>.
91. Meyersburg D, Welponer T, Kaiser A, Selhofer S, Tatarski R, Handisurya A, et al. Comparison of topical benzyl benzoate vs. oral ivermectin in treating scabies: A randomized study. *J Eur Acad Dermatol Venereol*. 2023;37:160-5, <http://dx.doi.org/10.1111/jdv.18573>.
92. Mytton O, McGready R, Lee S, Roberts C, Ashley E, Carrara V, et al. Safety of benzyl benzoate lotion and permethrin in pregnancy: A retrospective matched cohort study. *BJOG*. 2007;114:582-7, <http://dx.doi.org/10.1111/j.1471-0528.2007.01290.x>.
93. Gershanik J, Boecler B, Ensley H, McCloskey S, George W. The Gasping Syndrome and Benzyl Alcohol Poisoning. *N Engl J Med*. 1982;307:1384-8, <http://dx.doi.org/10.1056/NEJM198211253072206>.
94. Ko CJ, Elston DM. Pediculosis. *J Am Acad Dermatol*. 2004;50:1-12, [http://dx.doi.org/10.1016/S0190-9622\(03\)02729-4](http://dx.doi.org/10.1016/S0190-9622(03)02729-4).
95. Varela JA, Otero L, Espinosa E, Sánchez C, Junquera ML, Vázquez F. Phthirus pubis in a Sexually Transmitted Diseases Unit: A Study of 14 Years. *Sex Transm Dis*. 2003;30:292-6, <http://dx.doi.org/10.1097/00007435-200304000-00004>.
96. Burkhart CG, Burkhart CN. Oral ivermectin for Phthirus pubis. *J Am Acad Dermatol*. 2004;51:1037, <http://dx.doi.org/10.1016/j.jaad.2004.04.041>.
97. Bernigaud C, Fernando DD, Lu H, Taylor S, Hartel G, Chosidow O, et al. How to eliminate scabies parasites from fomites: A high-throughput ex vivo experimental study. *J Am Acad Dermatol*. 2020;83:241-5, <http://dx.doi.org/10.1016/j.jaad.2019.11.069>.