
GUIDELINES

Italian S3-Guideline on the treatment of Atopic Eczema - Part 1: Systemic therapy, adapted from EuroGuiDerm by the Italian Society of Dermatology and STD (SIDEMAST), the Italian Association of Hospital Dermatologists (ADOI) and the Italian Society of Allergological and Environmental Dermatology (SIDAPA)

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ABSTRACT

SIDeMaST (Società Italiana di Dermatologia Medica, Chirurgica, Estetica e delle Malattie Sessualmente Trasmesse) contributed to the development of the present guideline on the systemic treatment of chronic plaque psoriasis. With the permission of EuroGuiDerm, SIDeMaST adapted the guideline to the Italian healthcare context to supply a reliable and affordable tool to Italian physicians who take care of patients affected by atopic dermatitis. The evidence- and consensus-based guideline on atopic eczema was developed in accordance with the EuroGuiDerm Guideline and Consensus Statement Development Manual. Four consensus conferences were held between December 2020 and July 2021. Twenty-nine experts (including clinicians and patient representatives) from 12 European countries participated. This first part of the guideline includes general information on its scope and purpose, the health questions covered, target users and a methods section. It also provides guidance on which patients should be treated with systemic therapies, as well as recommendations and detailed information on each systemic drug. The systemic treatment options discussed in the guideline comprise conventional immunosuppressive drugs (azathioprine, ciclosporin, glucocorticosteroids, methotrexate and mycophenolate mofetil), biologics (dupilumab, lebrikizumab, nemolizumab, omalizumab and tralokinumab) and janus kinase inhibitors (abrocitinib, baricitinib and upadacitinib). Part two of the guideline will address avoidance of provocation factors, dietary interventions, immunotherapy, complementary medicine, educational interventions, occupational and psychodermatological aspects, patient perspective and considerations for pediatric, adolescent, pregnant and breastfeeding patients.

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KEY WORDS: Atopic dermatitis; Eczema; Guidelines.

Scoping and defining the purpose of the guideline

The aim of this guideline is to provide guidance on the management and treatment of patients with atopic ec-

zema (AE) of all severities and age groups. According to the scoping document, the objectives of the guideline are as follows:

- to generate recommendations and treatment algo-

ritms on topical therapy, phototherapy as well as novel and established systemic treatments for AE, based on the latest evidence;

- provide guidance in the management of AE patients during pregnancy and AE patients with allergic and other comorbidities.

Population and health questions covered by the guideline

The target population are patients with AE of all ages. Major health questions (regardless of sex, ethnicity or gender) regarding AE are as follows:

- what is the optimal treatment with regard to patients' needs, taking efficacy, safety/tolerability of different treatment options and comorbidities into consideration?;
- how should the selected treatment option best be managed and monitored?

Whenever possible and feasible, the recommendations are evidence-based, taking into account the results of systematic evidence synthesis based on rigorous methods and on the practical experience obtained by the expert group.

Targeted users of this guideline

This guideline has been prepared for physicians, especially dermatologists, pediatricians, allergists, general practitioners and other specialists taking care of patients with AE. Patients and caregivers may also be able to get reliable information and advice with regard to evidence-based therapeutic modalities.

Methods section

The EuroGuiDerm guideline on AE was developed in accordance with the EuroGuiDerm Methods Manual v1.3. For the detailed description of the guideline development process and an overview of the evidence referred to, please see the EuroGuiDerm guideline on AE Methods Report and the Evidence Report.¹

Both are available alongside the guideline document on the EDF website: <https://www.edf.one/de/home/Guidelines/EDF-EuroGuiDerm.html>.

Nomination of experts, management of conflict of interest

The guideline development group comprised 26 experts from 12 countries nominated by EuroGuiDerm national partner societies or the two guideline co-coordinators (A.W. and C.F.). All nominations were reviewed and confirmed by the

EuroGuiDerm Board of Directors. In addition, three patient representatives participated in the guideline development.

Thirty-eight percent of the experts declared personal-financial interests (for details on classification see EuroGuiDerm Methods Manual v1.3). *These members were neither eligible to take the lead in a respective working group nor for voting on recommendations pertaining to systemic treatment or on the stepped-care plan.*

Development of the guideline and the consensus process

The chapters of the guideline and the recommendations had been developed by the group members, who formed a number of working groups. Each chapter and all recommendations were reviewed, discussed and amended where appropriate by the entire group. All texts and recommendations were voted on with a necessary minimal agreement of >50% during the consensus conferences. AN facilitated all four consensus conferences using a structured consensus technique. Both internal and external review were conducted. Dissemination and implementation plans were developed. For more details, see Methods Report.

The wording of the recommendations was standardized (as suggested by the GRADE Working Group).²

Wording of recommendations

The recommendations are presented throughout this guideline as displayed below: alongside the wording of the recommendations (Table I, II) the arrow(s) and colors indicate the direction and the strength of each recommendation (Table III). The rate of agreement (consensus strength) is also displayed as the actual percentage and in form of a category-type pie chart. For all systemic drugs, we added the dosages (according to the European Medicines Agency).

Additionally, the certainty of evidence was added (Table III) where applicable (bold – significant difference; associations are reported in line with Drucker *et al.*).⁴

Evidence

The living systematic review by Drucker *et al.*⁵ was used as the evidence base based on which we created an evidence-to-decision framework (see Evidence Report). Furthermore, challenges exist with comparing clinical trials in AE due to their differences in trial design, including study comparators, rules for rescue treatment, washout periods for topical and systemic treatments, inclusion criteria and the duration of the screening period.⁶ Finally, this analysis does not take into consideration the overall management plan that targets long-term stabilization, flare prevention

TABLE I.—Terminology used throughout the guideline.

Terminology	Definition
Acute flare	Clinically significant worsening of signs and symptoms of AE requiring therapeutic intervention
Acute intervention	Treatments that address acute flares and typically lead to treatment response within days (in contrast to “maintenance treatment”)
Short term	When used in the context of clinical trials, this refers, to treatment up to 16 weeks
Reactive	Treatment initiations or adaptations in response to a visible change in disease severity, in particular disease flares (in contrast to “proactive treatment”)
Long term	When used in the context of clinical trials, this refers to treatment longer than 16 weeks
Proactive	Intermittent (typically twice a week) application of anti-inflammatory therapy to preciously affected skin, in addition to an ongoing emollient treatment of unaffected and affected skin (in contrast to “reactive treatment”)
Maintenance treatment	Regular, usually daily application of topical or systemic therapy for several months (in contrast to “acute intervention”)

TABLE II.—Definition of treatment goals.

Treatment goal	Definition
Remission/control	Satisfactory reduction in the signs and symptoms of AE whilst being on a safe long-term anti-inflammatory treatment
Complete remission	Disappearance of the signs and symptoms of AE without use of anti-inflammatory treatment

TABLE III.—Recommendation strengths: wording, symbols, interpretation and definition of certainty of evidence.³

Strength	Wording	Symbols	Implications
Strong recommendation for the use of an intervention	“We recommend...”	↑↑	We believe that all or almost all informed people would make this choice.
Weak recommendation for the use of an intervention	“We suggest...”	↑	We believe that most informed people would make this choice, but a substantial number would not.
No recommendation with respect to an intervention	“We cannot make a recommendation with respect to...”	0	At the moment, a recommendation in favor or against an intervention cannot be made due to certain reasons (e.g. no reliable evidence available, conflicting outcomes)
Weak recommendation against the use of an intervention	“We suggest against...”	↓	We believe that most informed people would make a choice against this intervention, but a substantial number would not.
Strong recommendation against the use of an intervention	“We recommend against...”	↓↓	We believe that all or almost all informed people would make a choice against this intervention.
High ⊕⊕⊕⊕	We are very confident that the true effect lies close to that of the estimate of the effect.		
Medium ⊕⊕⊕○	We are moderately confident in the effect estimate: the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.		
Low ⊕⊕○○	Our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect.		
Very low ⊕○○○	We have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of the effect.		

and avoidance of side-effects beyond 16 weeks.⁷ We only summarize the results here. For limitations, please refer to the website.

For each recommendation that is evidence-based, we added the certainty of the evidence when compared with placebo.⁴ The assessment of the certainty of evidence leads to four grades (Table 5.1. GRADE Handbook).³

Overview of recommendations

Stepped-care plans for the treatment of AE in adults (Figure 1) and in children and adolescents (Figure 2) can be found below. Table IV shows general recommendations for systemic drugs for adult AE patients, who are candidates for systemic treatment.

Introduction to the adaptation to the Italian Scenario of Healthcare

Introduction to adaptation to the Italian scenario of Healthcare and Regulations. The Italian Healthcare System (Servizio Sanitario Nazionale, SSN)

Italy has a public National Health Service, called “Servizio Sanitario Nazionale” (SSN) (Law 833 of 23 December 1978) with different regulatory and administrative competences at central, regional and local levels. The SSN offers the following services free of charges for everybody: routine medical examinations by a family doctor, medical examinations for acute conditions in an emergency room and hospitalization.

Examinations by a dermatologist or another specialist doctor have a copay up to 36.02 euros unless the patient has a low yearly income, particularly if he is elderly, or he is affected by a chronic and/or severe disease. However, AD is not in the list of diseases that give the right to exemption from the payment of the copay.

Examinations by a private doctor are not reimbursed by the SSN and a growing number of patients have a supplementary private insurance.

Citizens of EU countries can benefit from the Italian SSN through the European Health Insurance Card. Citizens of non-EU countries are assisted only for emergencies or acute conditions.

The Italian Medicines Agency (AIFA)

The Italian Medicines Agency (AIFA) intervenes in the governance of pharmaceutical expenditure by negotiating the price of medicinal products and managing the National Pharmaceutical Formulary.

In addition, AIFA monitors the appropriateness of use of medicines through the development, management and analysis of registers and consumption data.

It periodically reviews the Notes, which are regulatory tools that define the eligibility for reimbursement of some medicinal products and are updated on the basis of new scientific evidence and the needs of daily medical practice on the national territory.

Every month, the Agency also publishes the updated transparency lists, which are the lists of medicinal products whose patient coverage has expired, with the relative reference prices.

Introduction to systemic treatment

The area of systemic therapy of AE has flourished during the last few years, as many new substances are marketed, licensed, or in the last step of clinical development. The licensing programs of the various new biologics and small

molecules are providing much better levels of evidence than what is available for the longer existing drugs.

By tradition, systemic therapy of AE is deemed necessary if the signs and symptoms of AE cannot be controlled sufficiently with appropriate topical treatments and UV-light therapy. Systemic therapy can also be useful to reduce the total amount of topical corticosteroids (TCS) in patients who need large amounts of potent TCS for large body areas over prolonged periods to control their AE.

Candidates for systemic treatment may be either patients with a high composite score such as SCORAD above 50 (scale definition), or to patients clinically failing to respond to an appropriately conducted topical therapy (functional), or patients unable to participate in normal daily life activities whilst following an adequate treatment regimen (social definition).

Local regulations may necessitate the use of other scores such as physician-based scores (*e.g.* EASI) in combination with patient-reported outcomes (*e.g.* DLQI). Many other scores exist summarized and assessed by the HOME initiative that may also serve as a base to classify disease severity.⁸

It must be highlighted that the indication to systemic treatment is a patient individual decision, and that a signs-only score, such as EASI, is not an adequate tool to discriminate for providing or declining systemic therapy to an individual patient.

100% agreement.

Before starting systemic treatment, it is important to rule out relevant differential diagnoses such as cutaneous T-cell lymphoma and in selected cases primary immunodeficiency syndromes,⁹ and to ascertain that potential trigger factors such as allergic contact dermatitis, and behavioral as well as educational reasons for poor responses.

Until recently, rather broad acting immunosuppressants, such as systemic corticosteroids (SCS), ciclosporin (CyA), azathioprine (AZA), mycophenolate mofetil (MMF), enteric-coated mycophenolate sodium (EC-MPS) and methotrexate (MTX) were the only systemic treatment options available for difficult-to-treat AE. Most were not licensed for this indication (Table IV). These drugs may roughly be divided in two groups: SCS and CyA have a rapid onset of action and can be used to treat flares of AE or to bridge the time until onset of action of slow acting systemic immunosuppressants such as MTX, AZA and MMF/EC-MPS. The kinetics of the novel janus kinase inhibitors baricitinib (Bari), abrocitinib (Abro) and upadacitinib (Upa) place these

TABLE IV.—General recommendations for systemic drugs in adult AE patients who are candidate for systemic treatment (for details see corresponding chapter).

Recommendation	Conventional systemic treatments			Biologics		JAK-inhibitors		Rescue therapy
	Ciclosporin	Methotrexate	Azathioprine	Dupilumab	Tralokinumab	Baricitinib	Upadacitinib	Systemic corticosteroids
	↑↑	↑	↑	↑↑	↑↑	↑↑	↑↑	↑
Dose for adults ²	Licensed ≥16 years Standard dosage adults: 2.5-5 mg/kg per day in two single doses	Off-label; commonly used dosage adults: initial dose: 5-15 mg/week; max. dose: 25 mg/week	Off-label; commonly used dosage adults: 1-3 mg/kg per day	Licensed ≥6 years; adults: initially 600 mg s.c. day 1 followed by 300 mg Q2W	Licensed for adults; initially 600 mg sc. Day 1 followed by 300 mg Q2W; consider Q4W dosing at week 16 in those achieving clear or almost clear skin	Licensed for adults; dosage adults: 4 mg per day, reduction to 2 mg per day possible, depending on treatment response	Licensed ≥12 years; dosage adults: 15 or 30 mg per day based on individual patient presentation; age ≥65: 15 mg per day; the lowest effective dose for maintenance should be considered	General license for adults and children; dosage maximum: 1 mg/kg per day
Time to response (weeks) ⁴	1-2	8-12	8-12	4-8	4-8	1-2	1-2	1-2
Time to relapse (weeks, based on expert experience) ⁴	<2	>12	>12	>8	>8	<2	<2	<2
Monitoring	Complete blood count, renal and liver profile, blood pressure	Complete blood count, renal and liver profile, PIIINP in available, screen for chronic infections	Complete blood count, renal and liver profile, TPMT activity if available, screen for chronic infections	Not required	Not required	Complete blood count, lipid profile, liver profile	Complete blood count, lipid profile, liver profile	Not required for short-term treatment, consider blood glucose and testing for adrenal gland suppression with high doses/longer-term treatment
Selection of most relevant adverse events	Serum creatinine ↑, blood pressure ↑	Nausea, fatigue, liver enzymes ↑, myelotoxicity	Gastrointestinal disturbances, idiosyncratic hypersensitivity reactions, hepatotoxicity, myelotoxicity	Conjunctivitis, upper respiratory tract infections, arthralgia	Upper respiratory tract infections, conjunctivitis			Skin atrophy, weight gain, sleep disturbance, mood changes, hyperglycemia or new-onset diabetes, peptic ulcers/gastritis, osteoporosis

Symbols	Implications (adapted from GRADE) ²
↑↑	We believe that all or almost all informed people would make this choice
↑	We believe that most informed people would make this choice, but a substantial number would not
0	We cannot make a recommendation
↓	We believe that most informed people would make a choice against this intervention, but a substantial number would not
↓↓	We believe that all or almost all informed people would make a choice against this intervention
	No recommendation

¹SmPC; ⁴expert experience
 ↑: rise; AE: atopic eczema; GL: guideline; LDL: low-density lipoprotein; PIIINP: procollagen III N-terminal propeptide; TPMT: thiopurine-S-methyltransferase.

agents in the fast-acting group, whereas the Th2-blocking agents dupilumab (Dupi), tralokinumab and lebrikizumab, as well as the IL31-receptor blocking agent nemolizumab (Nemo) need some weeks to reach full efficacy.

Special considerations should be taken during the ongoing COVID-19 pandemic, as indicated by recommendations from the European Taskforce for Atopic Dermatitis.^{10, 11} Particular caution is required where patients receive combined systemic therapy.

The following recommendations for systemic drugs are based on expert opinions, the living systematic review by Drucker *et al.*,⁵ other published literature and medical considerations, and may differ from the legal licensing status and access routes, which are not uniform in European countries.

Classes of prescribability and reimbursement of drugs in Italy

Drugs in Italy have different classes of prescribability and reimbursement. Drugs for AD belongs to the A, C and H classes.

New expensive drugs, *e.g.* biologics and anti-JAK small molecules, are classified in the class H. The cost is fully paid in advance by the NHS and the drug is prescribed and distributed to the patients only by specialized hospital centers that are authorized by the regions.

All NHS doctors can prescribe class A drugs that are generally subsidized, requiring only a copay (the so called “ticket”). There are exemptions from the copay for specific categories of patients on the base of their annual income, age and pathological status. However, atopic dermatitis does not entitle to exemption.

If there are different products containing the same drug, only the cost of the cheapest product is fully reimbursed and the difference of the cost with the others should be paid by the patient. The price difference is particularly relevant if we compare the price of generic drugs with that of the original patented drugs.

If a class A drug is prescribed by a private doctor, the SSN does not reimburse the cost.

The cost of medicinal products in Class C is to be wholly borne by the customer.


Some medicines can be dispensed only following presentation of a medical prescription (range C with prescription), and other can be purchased directly without a medical prescription (over the counter, OTC, drugs), when the supervision of a doctor is not necessary for their use.

Online sales are permitted exclusively for OTC.

Emollients and other non-drug medications for atopic dermatitis are fully paid by patients, as well.

Conventional systemic drugs

Azathioprine (AZA)

<p>We suggest using azathioprine in AE patients who are candidates for systematic treatment.</p>	↑	<p>>75%</p>  <p>(15/16) Evidence and consensus based, see Evidence Report</p>
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Azathioprine: off-license; commonly used dosage:

Adults: 1-3 mg/kg per day
Children: 1-3 mg/kg per day

Certainty of evidence:^{4, 5}

Short term (8-16 weeks) vs. placebo (NMA main analysis)

⊕⊕⊕⊕ LOW for mean difference/standardized mean difference **change**

in signs, DLQI, Itch VAS; OR undesirable effects

⊕⊕⊕⊕ LOW for standardized mean difference **change in signs**, QoL

⊕⊕⊕⊕ VERY LOW for standardized mean difference change in itch

For azathioprine versus other drugs, see Evidence Report

Mechanisms of action and efficacy

AZA is a prodrug which is rapidly converted *in vivo* to the antimetabolite 6-mercaptopurine (6-MP), following cleavage of its imidazole side chain. It is believed to exert its primary immunosuppressant effect *via* metabolites of 6-MP, thioguanine nucleotides, which are subsequently incorporated into DNA, inhibiting its synthesis.¹²

The efficacy of AZA is comparable to that of MTX but lower compared to dupilumab and ciclosporin in clearing clinical signs of AE.⁴

Randomized clinical trials report a significant superiority of AZA vs placebo, with a decrease in clinical scores such as Six Area, Six Sign Atopic Dermatitis and Scoring Atopic Dermatitis (SASSAD) by 26% to 39% after 12 weeks.¹³ However, results from retrospective studies are less favourable with a percentage of AZA treatment failure varying from 30% to 57% due to adverse effects or lack of effectiveness.¹⁴⁻¹⁶ An observational follow-up study of 36 adult patients with severe AE treated with MTX or AZA over a 24-week period demonstrated less improvement in subjects with filaggrin mutations (36%, 13/36) compared with those without filaggrin mutations.¹³

Long-term studies on adult patients treated with either AZA or MTX showed a relative reduction in SCORAD of 53% (P<0.01) and 63% (P<0.01) after 2 years, and 54% and 53% after 5 years, respectively.^{13, 17} Patients with a Filaggrin mutation seemed to have slower but prolonged effects of therapy compared with patients without a mutation.^{13, 17}

Dosage: acute flare, short term, long term

- Off license;
- commonly used dosage:

- adults and children: 1-3 mg/kg bodyweight per day;
- If no improvement of AE occurs within 3 months, with- drawing azathioprine should be considered;
- we recommend combining AZA, as any systemic treatment with emollients and, whenever needed, topical anti- inflammatory treatment in AE patients;
- if timely thiopurine S-methyltransferase (TPMT) activity measurement is available, the following dosing of AZA has been suggested:
 - very low activity (<2.5 per mL red blood cells [RBC]), treatment should not be started;
 - intermediate activity (2.5-7.5 nmol/h/mL RBC): 0.5 mg/ kg bodyweight per day for the first 4 weeks and then increase to 1.0 mg/kg bodyweight per day;
 - normal activity (>7.5 nmol/h/mL RBC): 2.0 mg/ kg bodyweight per day for the first 4 weeks and then increase to 2.5-3.0 mg/kg bodyweight per day.

Low azathioprine doses (0.5-1.0 mg/kg bodyweight per day) for the first 4 weeks were shown to reduce gastrointestinal side effects.¹⁸

If TPMT results are not available prior to starting AZA therapy, then half the standard treatment should be given for about 4-6 weeks under close monitoring of full blood count and liver profile, prior to going up the full treatment dose.

Safety

In the short and medium term, the most commonly reported serious dose-dependent effects are hepatotoxicity and myelotoxicity, together with gastrointestinal disturbances. Furthermore, idiosyncratic hypersensitivity reactions (*e.g.* fever, rigors, myalgia, arthralgia and occasionally pancreatitis) may occur.¹⁹

Concerns have been raised about the potential carcinogenicity induced by long-term treatment with azathioprine (predominantly squamous cell skin cancer and non-Hodgkin's lymphoma), especially if AZA is combined with other immunosuppressant regimens.²⁰

Monitoring

- baseline: complete blood count, renal and liver profile;
- TPMT activity if available;
- screening for chronic infections (*e.g.* hepatitis B/C, HIV) before therapy should be considered;
- follow-up: Complete blood count, renal and liver profile twice monthly for 2 months, monthly for 4 months, then every other month and with dose increases;

- pregnancy testing before and during AZA therapy where indicated.

Combination with other treatments

Concomitantly to AZA, topical therapy with corticosteroids and or calcineurin inhibitors can be applied.

Because of a potentially increased risk to develop skin cancer, AZA should not be combined with UV light (UVA, UVB and PUVA).

Special considerations

There is a theoretical risk of teratogenesis with AZA. This is based on studies in animals in which very high doses of AZA were used. However, in practice, AZA has been used for over 30 years in sexually active men and women and no definite association between the drug and the incidence of fetal abnormalities has been observed. There also seems to be no effect on fertility.

According to a recent position paper by ETFAD,²¹ AZA use during pregnancy should be avoided as there are better options, but may be used off-label in the absence of other alternatives as continuation of treatment in women already receiving this treatment at the time of conception. According to experts' opinion of the ETFAD, the dosage of azathioprine should be reduced by 50% if it is continued during pregnancy. Initiation of azathioprine after conception is not recommend.

The use of AZA during lactation is debated. The WHO has recommended that the potential side-effects of AZA outweigh the effects and benefits of the treatment,²² and studies suggest that AZA intake during breastfeeding could increase the long- term risk of immunosuppression and carcinogenesis in the child.²³

AZA is not licensed for the treatment of AE in children but it has proven beneficial in several retrospective pediatric case series. The main disadvantage of AZA is that it reaches its maximum treatment effect only after 3-4 months.²⁴

Azathioprine, Methotrexate and Mycophenolate Mofetil

Azathioprine, Methotrexate and Mycophenolate Mofetil are not approved for the treatment of atopic dermatitis in Italy. Therefore, since the use if off-label, they are not reimbursed and the signing of an informed consent by patients is required. (<https://farmaci.agenziafarmaco.gov.it/bancadatifarmaci/>)

Stepped-care plan for adults with atopic eczema

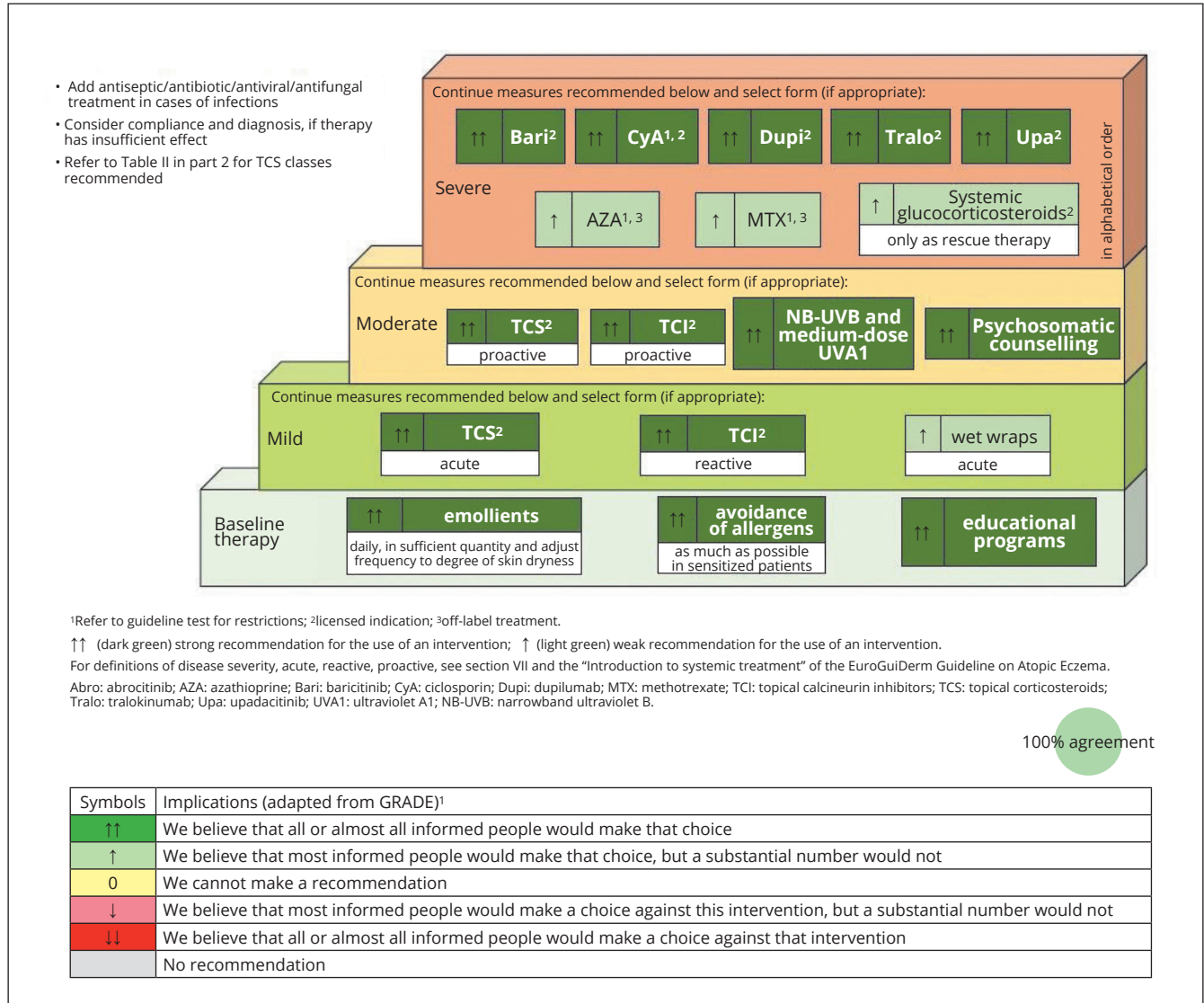


Figure 1.—Stepped-care plan for adults with AE.

Stepped-care plan for children and adolescents with atopic eczema

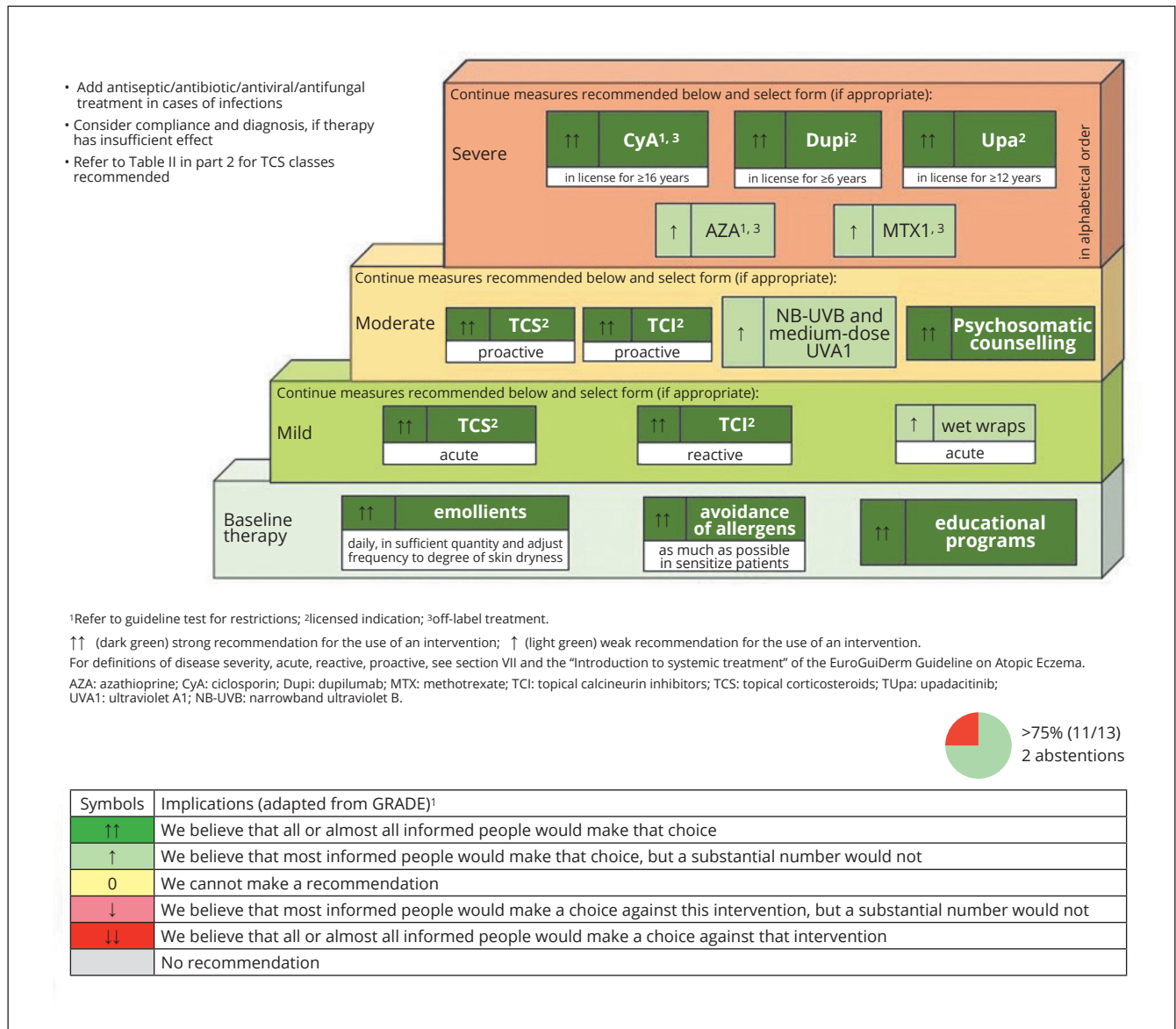





Figure 2.—Stepped-care plan for children and adolescents with AE.

Ciclosporin

We recommend using ciclosporin to achieve disease control in AE patients who are candidates for systemic treatment.	↑↑	>75%  (19/20) ² Evidence and consensus based, see Evidence Report
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Ciclosporin: in license for ≥16 years
 Standard dosage adults: 2.5-5 mg/kg per day in two single doses
 Commonly used dosage in children: 2.5-5 mg/kg per day in two single doses.
 Certainty of evidence:^{4, 5}
 Short term (8-16 weeks) vs. placebo (NMA analysis)
 ⊕⊕⊕⊕ LOW for mean difference/standardized mean difference **change in signs**, Itch VAS
 Short term (8-16 weeks) vs. placebo (NMA commonly used drugs)
 ⊕⊕⊕⊕ LOW for standardized mean difference **change in signs**, QoL
 ⊕⊕⊕⊕⊕ VERY LOW for standardized mean difference change in itch
 For ciclosporin versus other drugs, see Evidence Report
¹One abstention.

We recommend starting with higher ciclosporin dosage in order to achieve a more rapid response in AE patients who are candidates for systemic treatment.	↑↑	>75%  (16/17) Expert consensus
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We recommend close follow-up for potential blood pressure elevation and signs of renal impairment in AE patients on ciclosporin.	↑↑	>75%  (15/17) ² Expert consensus
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¹Two abstentions.

Mechanisms of action and efficacy

Ciclosporin inhibits T-cell activation and proliferation by blocking nuclear factor of activated T-cell (NFAT)-dependent cytokine production.

Ciclosporin has been approved for treatment of AE in adults in many European countries and is considered a first-line option for patients with severe disease if other, novel therapies are not available or indicated. Ciclosporin is very effective for AE in both children and adults with a better tolerability in children.^{25, 26} Although similarly effective in the above NMA meta-analysis evaluating trials up to 16 weeks, real-life data reveal a longer drug survival of dupilumab compared with CyA after 16 months.^{4, 27} In head-to-head trials ciclosporin was superior to MTX, prednisolone, IVIG, UVA and UVB, and similarly efficacious as EC-MPS.^{13, 28} In the short-term treatment of AE, higher ciclosporin dosages (5 mg/kg per day) lead to a more rapid response and are more efficacious than lower dosages (2.5-3 mg/kg per day).¹³ Long-term use of ciclosporin up to 1 year can be recommended based on several

trials; however, their evidence is limited because of the open-label design and high drop-out rates.¹³

Dosage: acute flare, short term, long term

- In license for ≥16 years;
- standard dosage adults: 2.5-5 mg/kg per day in two single doses;
 - acute flare, short-term: 4-5 mg/kg body weight per day;
 - long-term: 2.5-3 mg/kg body weight per day;
- commonly used dosage children: 2.5-5 mg/kg per day in two single doses;
 - we recommend combining CyA, as is the case with any systemic treatment, with emollients and, whenever needed, topical anti-inflammatory treatment in AE patients.

Safety

Ciclosporin has a narrow therapeutic index and requires a close follow-up for blood pressure and signs of renal impairment. Of note, clinically relevant increase of creatinine seems less common than expected.^{16, 26}

Monitoring

- Blood pressure, full blood count, renal and liver profile (including GGT) according to national guidelines (e.g. at base-line, 4 weeks and then 3-monthly);
- screening for hepatitis B/C and HIV before therapy should be considered.

Combination with other treatments

Concomitantly to ciclosporin, topical therapy with corticosteroids and/or calcineurin inhibitors can be applied.

Because of a potentially increased risk to develop skin cancer, ciclosporin should not be combined with UV light (UVA, UVB and PUVA).

Special considerations

Ciclosporin has been shown to be effective, safe and well tolerated in children and adolescents.^{25, 29}

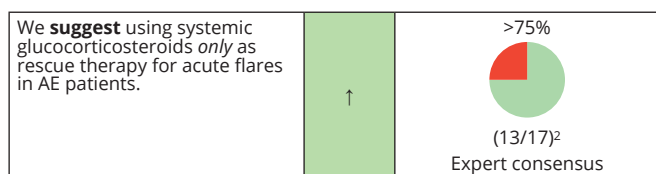
Ciclosporin can be considered in pregnant woman with severe AE. So far, no increased risk of congenital malformations or fetal death compared to the background populations have been reported. An increased risk of low birth-weight cannot be ruled out.²¹ Where systemic therapy is likely to be needed throughout pregnancy, ciclosporin is first-choice therapy.²¹

Oral cyclosporine is approved in Italy for the treatment of severe atopic dermatitis (criteria and/ or score of severity are not requested in detail). However, the use in adolescents younger than 16 years old is not recommended. Cyclosporine treatment should be prescribed by physicians experienced in the diagnosis and treatment of atopic dermatitis. Because of the variability of this disease, therapy must be individualized. The recommended dose range is 2.5-5 mg/kg/day in two divided oral doses. If an initial dose of 2.5 mg/kg/day does not result in a satisfactory response within 2-6 weeks of therapy, the daily dose can be rapidly increased to a maximum of 5 mg/kg. In very severe cases, rapid and adequate disease control is more likely to occur with an initial dose of 5 mg/kg/day. Once a satisfactory response has been achieved, the dose should be reduced gradually and, if possible, treatment with ciclosporin should be discontinued. A subsequent relapse can be treated with another course of cyclosporine. Although an 8-week course of therapy may be sufficient to achieve remission, treatment for up to 1 year has been shown to be effective and well tolerated provided monitoring guidelines are followed.

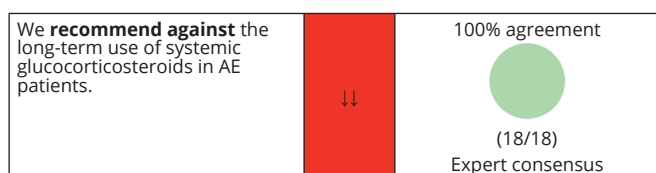
(https://farmaci.agenziafarmaco.gov.it/aifa/servlet/PdfDownloadServlet?pdfFileName=footer_000813_042787_RCP.pdf&sys=m0b113)

Only the cheapest generic product is fully reimbursed (https://www.gazzettaufficiale.it/atto/serie_generale/caricaDettaglioAtto/originario?atto.dataPubblicazioneGazzetta=2016-01-19&atto.codiceRedazionale=16A00318&elenco30giorni=false).

Systemic glucocorticosteroids



Systemic glucocorticosteroids: general unspecific license for steroid-responsive skin disease in adults and children; starting dose 0.5 mg/kg per day; dosage maximum: 1 mg/kg per day.
¹One abstention.



Mechanisms of action and efficacy

Glucocorticoids are a class of steroid hormones that bind to the glucocorticoid receptor. The activated glucocorticoid receptor complex upregulates the expression of anti-inflammatory proteins and suppresses the expression of pro-inflammatory proteins, leading to broad anti-inflammatory property.³⁰

There are only few studies in adult and pediatric AE patients, despite the regular use of systemic glucocorticosteroids in clinical practice. In studies conducted in children and adults, systemic glucocorticosteroids do not induce long-term remission and swift rebound is common. Systemic glucocorticosteroids have significantly inferior efficacy than ciclosporin.^{25, 31}

Dosage: acute flare, short term, long term

- Acute flare: starting dose is usually 0.5 mg/kg body-weight per day. Treatment should be discontinued or tapered as soon as possible;
- short-term and long-term: not recommended;
- we recommend combining systemic glucocorticosteroids, as is the case with any systemic treatment, with emollients and, whenever needed, topical anti-inflammatory treatment in AE patients.

Safety

Systemic glucocorticosteroids have a wide therapeutic index. Toxicity is related to the mean dose, cumulative dose and duration of use. At high doses and with long-term use (typically >0.5 mg/kg/day) important side-effects include skin atrophy, weight gain, sleep disturbance, mood changes, hyperglycemia or new onset diabetes, peptic ulcers/gastritis, osteoporosis, and increased susceptibility to infections.³² In particular with long-term use, patients can also develop adrenal suppression and together with a high risk of rebound flares when tapering the treatment dose, cessation can be challenging. Systemic glucocorticosteroids must therefore be avoided as a long-term treatment in adults and children. Even a fairly high dose can simply be stopped without tapering when used for no longer than 3 weeks.³³

Monitoring

For acute rescue therapy, there is no standard set of laboratory parameters. Monitoring should there be based on individual patient needs.

Combination with other treatments

None of the other treatments in AE are contraindicated when using systemic glucocorticosteroids.


Special considerations

Treatment of acute flares of AE with oral glucocorticosteroids is moderately effective.^{25, 31}

Systemic glucocorticosteroids have an unfavourable risk/benefit ratio for the long-term treatment of adult and pediatric AE.

Systemic corticosteroids are approved in Italy for the treatment of severe or debilitating allergic conditions, unresponsive to other therapies, such as bronchial asthma, contact dermatitis and atopic dermatitis (https://farmaci.agenziafarmaco.gov.it/aifa/servlet/PdfDownloadServlet?pdfFileName=footer_000794_010089_RCP.pdf&retry=0&sys=m0b113).

Methotrexate

We suggest using methotrexate in AE patients who are candidates for systemic treatment.	↑	100% agreement  (17/17) Evidence and consensus based, see Evidence Report
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Methotrexate: off license; commonly used dosage
 adults: initial dose: 5-15 mg per week; maximum dose: 25 mg per week
 children: 0.3-0.4 mg/kg per week; maximum dose: 25 mg per week
 Certainty of evidence:^{4, 5}
 Short term (8-16 weeks) vs. placebo (NMA main analysis)
 ⊕⊕○○ LOW for standardized mean difference **change in signs**, QoL
 Short term (8-16 weeks) vs. placebo (NMA currently used drugs)
 ⊕⊕○○ LOW for standardized mean difference **change in signs**, QoL
 ⊕○○○ VERY LOW for standardized mean difference change in itch
 For methotrexate versus other drugs, see Evidence Report

Mechanisms of action and efficacy

MTX is a folic acid antagonist that impedes cell division, DNA/RNA synthesis and repair and protein synthesis, altogether suppressing the activity of the immune system. Although its exact action in AE is not fully understood, inhibition of the janus kinase (JAK)/STAT pathway has been proposed.³⁴

MTX has been used in the treatment of moderate and severe AE for years, but only a limited number of non-randomized controlled trials have examined the effect and treatment regimens. Consequently, recommendations have been primarily based on case series and expert consensus,³⁵⁻³⁷ one controlled study comparing MTX with AZA in adults³⁸ and an open-label randomized multicenter study in children.³⁹ Altogether these studies support that MTX can be considered a moderately effective, relatively safe and well-tolerated treatment for severe AE both in children and adults – findings also in keeping with recent retrospective studies.⁴⁰⁻⁴² The efficacy of MTX was comparable to AZA and lower

than dupilumab and ciclosporin in clearing clinical signs of AE at week 16. However, there are no long-term follow-up head-to-head studies available for further comparison.⁴ The onset of action takes several weeks and peak efficacy is seen after months, but speed of treatment effect onset depends on the dosing regimen.³⁵⁻³⁷ One adult study suggests that patients who do not benefit from a moderate weekly dose (10-15 mg) of MTX over a 3-month treatment period will probably not benefit from an increased dosage. However, slow gradual up-dosing of MTX might underestimate the therapeutic potential of the drug in AE. In children 0.4 mg/kg/week is recommended, which is significantly higher than dosing in adults.³⁵ 25 mg per week is the widely used maximum treatment dose for adult and pediatric AE patients.

Dosage: acute flare, short term, long term

- Off-license;
- commonly used dosage;
 - adults: initial dose: 5-15 mg/ per week; maximum dose: 25 mg/week;
 - children: 0.3-0.4 mg/kg per week
- Acute flare and short-term: no relevant dosing;
 - oral and subcutaneous delivery are considered equivalent options of administration. For patients in whom MTX 15 to 25 mg orally once weekly is ineffective or poorly tolerated, a trial of subcutaneous MTX administration is an alternative;
 - we recommend combining MTX, as is the case with any systemic treatment, with emollients and, whenever needed, topical anti-inflammatory treatment in AE patients;
 - concomitant use of folic acid should be considered to reduce gastrointestinal and other side-effects related to the folic acid antagonistic effect of the drug.⁴³

Safety

As MTX is a commonly used drug in dermatology, the safety profile is well recognized, with nausea, fatigue and raised liver enzymes as main side-effects, while pancytopenia and idiopathic pulmonary fibrosis are of key concern but only very rarely seen.

MTX is generally well tolerated and is considered safe for long-term treatment, based on experience and multiple studies including both adults and children suffering from psoriasis and rheumatologic disease.^{44, 45}

Monitoring

Complete blood count, renal and liver profile before and every 4 weeks for the first 3 months or, after increasing the dose, then every 8-12 weeks.

Type III procollagen peptide (PIIINP) should be monitored according to national and local guidelines when available. Fibrosan or liver biopsy when necessary in selected cases.

Screening for chronic infections (e.g. hepatitis B/C, HIV, tuberculosis) before therapy should be considered.

Any noteworthy impact on liver or bone marrow function should give cause to dose reduction or transient or total discontinuation of treatment.

Combination with other treatments

Combination with TCS, TCI or narrow band UV phototherapy are established treatment combinations and considered safe. Concomitant use of ciclosporin is a relative contraindication. There is experience from rheumatoid arthritis for combining with the JAK inhibitor baricitinib.

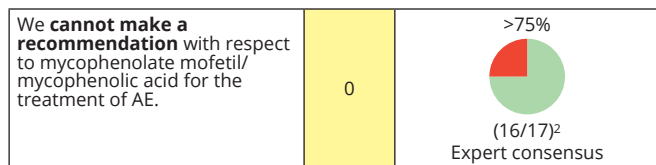
Special considerations

MTX may be used for treatment of AE in both adults and children.

Subcutaneous administration increases bioavailability and tolerability, as well as adherence, compared to oral treatment.

MTX affects fertility and is teratogenic. Fertile women should use effective contraception. The same is recommended for men treated with MTX living with a woman of childbearing potential.

Mycophenolate mofetil



Mycophenolate mofetil: off-license; commonly used dosage
 adults: 1-3 g per day
 children: 30-50 mg/kg per day
¹One abstention.

Mechanisms of action and efficacy

Mycophenolate mofetil is a prodrug of mycophenolic acid (MPA), an inhibitor of inosine-5⁰-monophosphate dehydrogenase. MPA depletes guanosine nucleotides preferentially in T and B lymphocytes and inhibits their proliferation. MPA also inhibits the glycosylation and expression of adhesion molecules, and the recruitment of lymphocytes and monocytes into sites of inflammation.⁴⁶

A recent systematic review and meta-analysis⁴⁷ including 18 studies with a total of 140 adult and pediatric patients evaluated the efficacy of off-label use of MMF in patients with AE refractory or not tolerating other first-line systemic agents. There was a significant reduction in pre- to post-SCORAD scores by 18 points (P=0.0002) with 77.6% of patients reporting partial or full remission. Relapses occurred in 8.2% of cases. The average time for initial effects was 6.8 7 weeks.

Dosage: acute flare, short term, long term

- Off-license;
- commonly used dosage:
 - adults: 1-3 g per day;
 - children: 30-50 mg/kg bodyweight per day;
 - typically given in two divided doses;
- we recommend combining MMF, as is the case with any systemic treatment, with emollients and, whenever needed, topical anti-inflammatory treatment in AE patients.

Safety

The most common side-effects include headaches and gastrointestinal symptoms, followed by infections, especially during long-term therapy.

Hematological adverse effects include anemia, leukopenia, neutropenia and thrombocytopenia, albeit rarely.

Monitoring

- Complete blood count, renal and liver profile before therapy, then every 2 weeks for 1 month; monthly for 3 months; every 2-3 months thereafter;
- screening for chronic infections (e.g. hepatitis B/C, HIV) according to national and local guidelines;
- pregnancy testing before and during MMF therapy if indicated.

Combination with other treatments

Concomitantly to MMF, topical therapy with corticosteroids and/or calcineurin inhibitors can be applied.

Special considerations

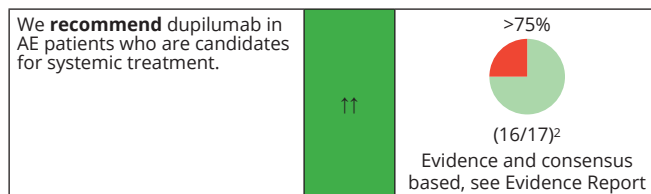
In case series, the efficacy and safety of MMF in children have been investigated. The drug has shown a positive treatment response with minimal adverse effects and appears to be better tolerated than AZA.⁴⁸

Biologics

Regulations for prescribability and reimbursement of all biologics and small molecules for AD in Italy

All biologics and small molecules for AD are class H drugs *i.e.* they are fully reimbursed by the Italian NHS only if they are prescribed by dermatologists and hospital centers that are selected by the Health Departments of the Regions. Patients must be included in dedicated AIFA Monitoring web-based registries.

Dupilumab



Dupilumab: in license for ≥ 6 years;
 Age 6-11: from 15 kg <60 kg, initially 300 mg s.c. day 1 and 15 followed by 300 mg Q4Wm when ≥ 60 kg, initially 600 mg s.c. day 1 followed by 300 mg Q2W

Age 12-17: <60 kg: initially 400 mg s.c. day 1 followed by 200 mg Q2W, when ≥ 60 kg: initially 600 mg s.c. day 1 followed by 300 mg Q2W
 Adults: initially 600 mg s.c. day 1 followed by 300 mg Q2W

Certainty of evidence:^{4,5}
 Short term (8-16 weeks) vs. placebo (NMA main analysis)
 ⊕⊕⊕⊕ HIGH for mean difference/standardized mean difference **EASI, change in signs, POEM, DLQI**
 ⊕⊕⊕○ MODERATE for undesirable effects
 Short term (8-16 weeks) vs. placebo (NMA currently used drugs)
 ⊕⊕⊕⊕ HIGH for standardized mean difference **change in signs, QoL, change in itch**
 Long term (52 weeks) vs. placebo
 RoB low for change in **EASI, POEM, DLQI, undesirable effects**
 For dupilumab versus other drugs, see Evidence Report
¹One abstention.

Mechanisms of action and efficacy

Dupilumab is the first marketed fully human IgG4 monoclonal antibody (mAb) in the treatment of AE and has been available for treatment of adults for more than 2 years in many countries. Recently, it has also been approved for adolescents and children from 6 years of age in some countries. Dupilumab binds to the α -subunit of the IL-4 receptor, which is part of both the IL-4 and the IL-13 receptor complex. The safety and efficacy of dupilumab was primarily established in placebo-controlled studies in moderate-to-severe AE.⁴⁹ Dupilumab showed significant clinical effects across 3 distinct severity assessment tools: Eczema Area and Severity Index (EASI), Investigator's Global Assessment (IGA) and SCORing Atopic Dermatitis (SCORAD). Moreover, dupilumab treatment significantly reduced pruritus. Dupilumab has shown efficacy in both intrinsic and extrinsic AE.⁵⁰ Dupilumab is also registered for treatment of moderate-to-severe asthma,

eosinophilic esophagitis, and chronic rhinosinusitis with nasal polyps, thereby covering several type 2 inflammatory diseases.

Dosage: acute flare, short term, long term

The approved dosing of dupilumab in adults consists of a 600 mg subcutaneous loading dose followed by maintenance doses of 300 mg every other week (Q2W). For children, the following dosing regimens are used: licensed for ≥ 6 years; age 6-11: from 15 kg <60 kg, initially 300 mg s.c. day 1 and 15 followed by 300 mg Q4W, when ≥ 60 kg, initially 600 mg s.c. day 1 followed by 300 mg Q2W age 12-17: <60 kg: initially 400 mg s.c. day 1 followed by 200 mg Q2W, when ≥ 60 kg: initially 600 mg s.c. day 1 followed by 300 mg Q2W.

Dupilumab has been used in an open-label study for up to 3 years in adults with moderate-to-severe AE, but some former trial patients have continued open label on the medication much longer. Safety data were consistent with previously reported trials and the known dupilumab safety profile.⁵¹

Safety

Dupilumab treatment is in generally well tolerated, and routine blood tests are not recommended, but a substantial number of patients develop conjunctivitis (over 30% in some 'real world' settings), of which most are mild-to-moderate.^{52, 53} Topical treatment with anti-inflammatory eyedrops is often sufficient, without need to discontinue treatment.⁵⁴

Monitoring

No biochemicals or instrumental exams are reported to be required for the monitoring of the therapy.

Combination with other treatments

An additional phase III trial, evaluated dupilumab treatment and a concomitant topical corticosteroid (TCS) compared with placebo and a concomitant TCS over 52 weeks.⁵⁵ The co-primary end points included IGA score of 0 or 1 and EASI-75, were assessed at week 16: more patients who received dupilumab plus topical corticosteroids achieved the co-primary endpoints of IGA 0/1 and EASI 75. Results at 52 weeks were similar. Approximately 15% more subjects achieved a 75% reduction in the EASI score at week 16 in this trial compared with previous phase III studies where dupilumab was administered as monotherapy.⁴⁹

Combination therapy with TCS, TCI, and UV light treatment is well established.

Special considerations

AE patients with type 2 comorbidities such as asthma, allergic rhino-conjunctivitis with nasal polyps, or eosinophil esophagitis may also have beneficial effects of dupilumab treatment on these diseases.

The approved indications of Dupilumab (Dupixent®, Sanofi-Aventis Groupe, Paris, France) in Italy are:

Treatment of moderate to severe atopic dermatitis in adults eligible for systemic therapy with EASI score ≥ 24 and for whom ciclosporin treatment is contraindicated, ineffective or not tolerated), (Determina n. DG/133/2018, GU n.208 del 7-9-2018). Adolescents aged 12 years or over (Determina n. DG/1203/2020 GU n. 305 del 09-12-2020) and children aged 6 to 11 years (Determina n.115/2022, GU n.42 del 19-02-2022) can be treated with Dupilumab if EASI score ≥ 24 or they present one of the following characteristics: localization in visible and/or sensitive areas; NRS scale of pruritus ≥ 7 ; quality of life with DLQI index ≥ 10)

Lebrikizumab

Lebrikizumab is currently not licensed for any indication world- wide. Therefore, we do not give a specific recommendation for its use in AE.

Mechanisms of action and efficacy

Lebrikizumab is a high- affinity humanized immunoglobulin G4 mAb that binds specifically to soluble interleukin 13 and selectively prevents formation of the IL-13Ra1/IL-4Ra heterodimer receptor signaling complex. In a randomized, placebo-controlled, double-blind, phase IIb study, adults with moderate-to-severe AE patients were randomized to placebo every 2 weeks or to subcutaneous injections of lebrikizumab at the following doses: 125 mg every 4 weeks (250 mg loading dose [LD]), 250 mg every 4 weeks (500-mg LD), or 250 mg every 2 weeks (500 mg LD at baseline and week 2).⁵⁶

Compared with placebo, lebrikizumab groups showed dose- dependent, statistically significant improvement in EASI scores, pruritus NRS score, POEM and IGA.⁵⁶

Dosage: acute flare, short term, long term

Although all the different dosages of lebrikizumab proved to be effective, optimal dosing regimens have yet to be determined. Phase 3 studies are currently underway

testing lebrikizumab 250 mg Q2W the induction phase, and both 250 mg Q2W and Q4W in the maintenance phase.

Safety

Treatment-emergent adverse events were reported in 24 of 52 placebo patients (46.2%) and in lebrikizumab patients as follows: 42 of 73 (57.5%) for 125 mg every 4 weeks, 39 of 80 (48.8%) for 250 mg every 4 weeks, and 46 of 75 (61.3%) for 250 mg every 2 weeks; most were mild-to-moderate and did not lead to discontinuation. In all lebrikizumab groups, herpes virus infections and conjunctivitis were reported at low rates.

Simpson *et al.* reported injection site reactions (1.3%), herpes infection (3.8%), eosinophilia (3.2%) with no associated clinical symptoms, and conjunctivitis (9.6%) as adverse events in patients treated with lebrikizumab.⁵⁷

Notably, lebrikizumab appears to have lower rates of ocular complications than dupilumab.

Monitoring

No biochemical or instrumental examinations are reported to be required for the monitoring of the therapy.

Combination with other treatments

The use of topical corticosteroids during the flares of AE could be useful in combination with lebrikizumab, and is under investigation in the phase 3 program.

At present, the EMA has accepted the filing of the Marketing Authorization Application (MAA) for lebrikizumab for the treatment of moderate to severe atopic dermatitis and an approval is pending. The documentation will then be sent to AIFA.

Nemolizumab

Nemolizumab is currently not licensed for any indication world- wide. Therefore, we do not give a specific recommendation for its use in AE.

Mechanisms of action and efficacy

Nemolizumab is a humanized mAb targeting the IL-31 receptor alpha chain (IL-31RA), which was initially developed for the treatment of AE-related pruritus.

In a phase II, randomized, double-blind, placebo-controlled, 12-week trial, nemolizumab at monthly doses significantly improved pruritus.⁵⁸

In a 2b study with nemolizumab 30 mg dosing and TCS, there were significant improvements in signs and

symptoms of AE - EASI scores, PP-NRS, sleep and DLQI score, which was confirmed in a *post-hoc* sub-analysis of the EASI ≥ 16 cohort.^{59, 60}

In a recently published 16-week, double-blind, phase III trial, Japanese patients with AE and moderate-to-severe pruritus received subcutaneous nemolizumab (60 mg) or placebo every 4 weeks until week 16, with concomitant topical agents.⁶¹ The primary end point was the mean per cent change in the visual-analogue scale (VAS) score for pruritus from baseline to week 16. Secondary end points included the time course of change in the VAS score for pruritus up to week 4, EASI score, DLQI, Insomnia Severity Index, and safety. At week 16, the mean per cent change in the VAS score was -42.8% in the nemolizumab group and -21.4% in the placebo group. The use of subcutaneous nemolizumab in addition to topical agents for atopic dermatitis resulted in a highly significant reduction in pruritus than placebo plus topical agents.

Dosage: acute flare, short term, long term

The first phase II study investigating nemolizumab published in 2017 investigated 0.1, 0.5, 2 mg/kg dosages administered every 4 weeks and 2 mg/kg dosage administered every 8 weeks. Results at 12 weeks found a significant, dose-dependent improvement in the primary outcome of pruritus for all groups that received nemolizumab every 4 weeks, as compared with placebo.⁵⁸ In a two-part, phase II, randomized control trial published in 2018, Kabashima *et al.*⁶¹ compared three different nemolizumab dosages: 0.1, 0.5, 2 mg/kg administered every 4 weeks and 2 mg/kg administered every 8 weeks. All the parameters considered in the study showed an improvement, and no evidence was found that the highest dosage was more effective than the lowest. Furthermore, the study showed that the positive outcomes obtained with nemolizumab were maintained for up to 64 weeks.

In another 24-week, randomized, double-blind, multicenter study published in 2019 by Silverberg *et al.*,⁵⁹ three different nemolizumab dosages, 10, 30 and 90 mg, were compared in an ethnically more diverse population. The drug was administered once every 4 weeks and nemolizumab 30 mg showed maximum dosage efficacy in improving EASI, IGA, and pruritus.

In the latest published study conducted in Japanese patients,⁶¹ the dosage tested was 60 mg, administered every 4 weeks. At the reported dosage, nemolizumab showed a greater efficacy in reducing pruritus, compared to placebo plus topicals.

Safety

The most frequent adverse events related to the drug are reported to be injection-related reactions, musculoskeletal and connective tissue symptoms, upper respiratory tract infections, nasopharyngitis, peripheral oedema and increased creatine phosphokinase.⁵⁹

The authors conclude that longer and larger trials are necessary to determine whether nemolizumab has a durable effect and is safe for AE patients.⁵⁹

Monitoring


No biochemical or instrumental exams are reported to be required for the monitoring of the therapy.

Combination with other treatments

According to the available study trials, the use of topical treatments such as emollients, corticosteroids and calcineurin inhibitors as a rescue therapy, in addition to nemolizumab, could have a synergistic effect in the treatment of AE and AE-related pruritus.

EMA has granted a product specific waiver for nemolizumab for the indication “prurigo nodularis” (20 March 2020). At present, no approval request is pending at EMA and AIFA for the indication “atopic dermatitis”.

Omalizumab

<p>We cannot make a recommendation with respect to the use of omalizumab for the treatment of AE.</p>	<p>0</p>	<p>100% agreement  (17/17) Expert consensus</p>
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Omalizumab: in label for allergic asthma (≥ 6 years), chronic rhinosinusitis with nasal polyps (CRSwNP) (≥ 18 years) and chronic spontaneous urticaria (≥ 12 years).

Commonly used dosage:

Dosage (allergic asthma and CRSwNP): depends on baseline IgE (IU/mL), measured before the start of treatment, and body weight. The maximum recommended dose is 600 mg omalizumab every two weeks. Please refer to the SmPC for further details. Dosage (chronic spontaneous urticaria): 300 mg every four weeks.

Mechanisms of action and efficacy

Most AE patients have elevated serum IgE levels, but the pathogenic role of IgE in AE remains unknown. The anti-IgE antibody omalizumab has been used with great suc-

cess for treatment of chronic spontaneous urticaria (CSU). A recent systematic review and meta-analysis has assessed the preclinical and trial data regarding omalizumab treatment of AE, which are conflicting.⁶²

Omalizumab is licensed for treatment of asthma and CSU, but not for treatment for AE.

Omalizumab binds free IgE, which leads to immune complexes of IgE and omalizumab. IgE bound to omalizumab cannot bind to the alpha chain of the high-affinity receptor for IgE, thereby inhibiting its binding to mast cells, basophils and epidermal dendritic cells,^{63, 64} and subsequent immunological effects.

There are many case reports and case series,⁶² but only few controlled trials studying omalizumab treatment of AE.^{62, 65} In summary, the data show a measurable, but moderate efficacy of omalizumab for improving signs and symptoms of AE.^{62, 66} There is no predictive marker linked to a better clinical response, and most of the published evidence is of low quality. The safety of omalizumab is very good,⁶² but the unpredictable and statistically low efficacy prevents a general recommendation for omalizumab regarding treatment of AE.

Dosage: acute flare, short term, long term

ADULT

Different dosages have been tested in AE patients, ranging from 150 to 450 mg every 2 weeks or every 4 weeks. A recent systematic review and meta-analysis by Wollenberg *et al.* found that patients with lower baseline IgE showed a positive response to treatment with omalizumab compared with patients with very high-to-extremely high serum IgE.⁶²

An older systematic review and meta-analysis by Wang *et al.* also found that IgE serum concentrations of less than 700 IU/mL were associated with a better clinical response, compared with with IgE concentrations of 700 to >5000 IU/mL. Age, sex, baseline clinical disease severity, a history of concomitant asthma, and the use of 600 mg/month or more of omalizumab showed no significant association with the clinical results associated with omalizumab use.⁶⁷

CHILDREN

The ADAPT (Atopic Dermatitis Anti-IgE Paediatric Trial) trial evaluated the possible role of omalizumab in the management of severe pediatric AE with concomitant allergic disease (asthma, allergic rhino-conjunctivitis or food allergies) for 24 weeks. The drug dose was determined by baseline total IgE (range: 30 to 1500 IU/mL), measured

before the start of treatment, and body weight (kg) and calculated using the formula: 0.0169 weight (kg) x total IgE level (kU/L) in 2-4 weekly injections. The study showed that omalizumab significantly reduced disease severity and improved QoL in pediatric patients with severe AE and highly elevated IgE levels (median baseline total IgE of 8373 IU/L) compared with placebo.⁶⁵ However, this improvement was below the minimal clinically important difference for the main outcome (objective SCORAD).

Safety

There is a general consensus about the overall good safety profile of omalizumab, with some controlled studies reporting excellent tolerability up to 4 years. A 2009 revision of data from controlled trials concluded that incidence of anaphylaxis was 0.14% in omalizumab-treated patients and 0.07% in control subjects. Of note, no serum-sickness attributable to the drug and no anti-omalizumab antibodies have been reported to date.⁶⁸

There are no reported interactions of omalizumab with other medications used for AE or other allergic diseases. If clinically needed, omalizumab may be considered during pregnancy. More attention has been put over the appearance of gut parasite infections in treated patients, as IgE is an important player in the host defense against parasitic helminths. A randomized placebo- controlled trial in 137 adult subjects with respiratory allergy at high risk of helminth infection showed a modest increase of the incidence of parasitism in the active group.⁶⁹

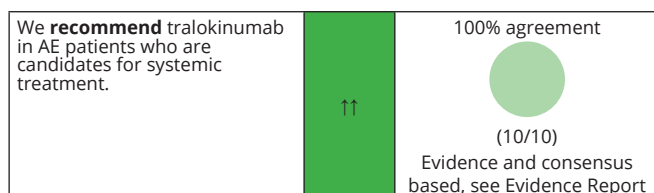
Monitoring

No biochemicals or instrumental examinations are reported to be required for the monitoring of the therapy. IgE levels increase following administration of omalizumab and may remain elevated for up to 1 year following discontinuation of the drug.

Omalizumab is not approved for the treatment of atopic dermatitis in Italy. It is approved for adults, adolescents and children aged between 6 and <12 years with established IgE (immunoglobulin E) mediated asthma; adults and adolescents (12 years of age and older) and children (6 to <12 years of age), as an adjunctive therapy, to improve the asthma control in patients with persistent severe allergic asthma which were positive in the skin test or *in vitro* reactivity to a perennial aeroallergen and have frequent daytime symptoms or nocturnal awakenings and in patients with documented repeated severe asthma exacerbations, despite daily intake of high doses of corticosteroids

plus a long-acting beta2-agonist by inhalation. In addition, it is indicated, as adjunctive therapy, for the treatment of chronic spontaneous urticaria in adult and adolescent patients (12 years of age or older) with inadequate response to treatment with oral H1 antihistamines. The prescribability is acknowledged by the Regional Health Departments to specialists of selected hospitals and a therapeutic plan must be filled in a dedicated web-based registry (https://www.aifa.gov.it/documents/20142/961234/Determina_DG-1111-2021_Xolair.pdf)

Tralokinumab



Tralokinumab: in license adults; Short term (8-16 weeks) vs. placebo (NMA main analysis) dosage adults: initially 600 mg s.c. day 1 followed by 300 mg Q2W at prescriber's discretion, every fourth week dosing may be considered for patients who achieve clear or almost clear skin after 16 weeks of treatment. Certainty of evidence:^{4,5} ⊕⊕⊕⊕ MODERATE for mean difference/standardized mean difference EASI, DLQI ⊕⊕⊕⊕ LOW for undesirable effects For tralokinumab versus other drugs, see Evidence Report

Mechanisms of action and efficacy

Tralokinumab is a fully human, high-affinity IgG4 mAb, which neutralizes IL-13, and has been approved by the EMA in summer 2021.⁷⁰ In two 52-week, double-blind, placebo-controlled phase III trials, adults with moderate-to-severe AE were randomized to subcutaneous tralokinumab 300 mg every 2 weeks or placebo.⁷¹ Tralokinumab monotherapy was superior to placebo at 16 weeks of treatment. Coprimary end points were IGA score of 0 or 1 and EASI 75 at week 16. Patient achieving an IGA score of 0/1 and/or EASI 75 with tralokinumab at week 16 was rerandomized to tralokinumab Q2W or every 4 weeks or placebo for 36 weeks. The majority of week 16 tralokinumab-responders maintained response at week 52 with continued tralokinumab treatment without any rescue medication.

Dosage: acute flare, short term, long term

The recommended dosage is 300 mg every 2 weeks after a loading dose of 600 mg at treatment onset. At prescriber's discretion, every fourth week dosing may be considered for patients who achieve clear or almost clear skin after 16 weeks of treatment.

Phase III trials have also investigated what happens

when patients who do well for 16 weeks on tralokinumab continue treatment as labelled, reduce treatment frequency, or discontinue treatment.

After 16 weeks, patients who reached EASI 75 or IGA success were rerandomized to continue treatment every 2 weeks, titrate down to every 4 weeks, or use placebo. At 52 weeks, without TCS, more than 55% of patients who continued twice-monthly treatment maintained EASI 75, as did approximately 50% of patients treated monthly. More than 51% of patients who stayed on twice-monthly dosing maintained IGA 0 or 1, vs. 39% and 45% of patients who switched to monthly dosing.

Safety

In the two studies, adverse events were reported in 76.4% and 61.5% of patients receiving tralokinumab and in 77.0% and 66.0% of patients receiving placebo in the 16-week initial period.

Notably, tralokinumab appears to have lower rates of ocular complications than dupilumab.⁷¹

The combination therapy with TCS, TCI and UV light treatment is possible.

Monitoring

No biochemical or instrumental examinations are reported to be required for the monitoring of the therapy.

Combination with other treatments

In an additional phase III double-blind, placebo study the efficacy and safety of tralokinumab in combination with TCS as needed in patients with moderate-to-severe AE were evaluated. At week 16, significantly more tralokinumab-treated patients than placebo achieved IGA 0/1 and EASI 75. Nine of ten EASI 75 responders at week 16 maintained response at week 32 with continued tralokinumab and TCS as needed.⁷²

The approved indication of Tralokinumab (Adtralza, Leo Pharma, Ballerup, DK) in Italy is the treatment of adolescents 12 years of age and older and adult patients with severe atopic dermatitis (EASI ≥ 24). https://www.gazzettaufficiale.it/atto/serie_generale/caricaDettaglioAtto/originario?atto.dataPubblicazioneGazzetta=2022-0804&atto.codiceRedazionale=22A04366&elenco30giorni

The recommended initial dose is 600 mg (four 150 mg subcutaneous injections) followed by 300 mg (two 150 mg injections) at two weeks' intervals.

Intervals of four weeks may be considered for patients with clear or nearly clear skin after 16 weeks of treatment. The

discontinuation of treatment should be evaluated in patients without response after 16 weeks of treatment. Some patients with initial partial response may subsequently improve further with continued treatment every two weeks beyond 16 weeks.

Tralokinumab can be used with or without topical corticosteroids. The use of topical corticosteroids, if appropriate, may provide an additive effect to the overall efficacy of tralokinumab. Topical calcineurin inhibitors can be used, but should be reserved only for problem areas, such as the face, neck, intertriginous areas and genitals. (<https://www.codifa.it/farmaci/a/adtralza-tralokinumab-dermatologici>)

JAK-inhibitors

The JAK family, constituting JAK1, JAK2, JAK3 and tyrosine kinase 2 (TYK2), are a class of cytoplasmic tyrosine kinases.⁷³ JAKs dock to the intracellular part of cytokine receptor chains to generate functional signaling complexes and regulate the inflammatory process through activating the intracytoplasmic transcription factors termed as signal transducer and activator of transcription (STAT). When activated, STAT proteins produce dimers, which translocate into the nucleus and either positively or negatively regulate downstream target gene expression of inflammatory mediators, suggesting that inhibiting JAK activity may be more effective than targeting a single cytokine. Beyond the disruption of cutaneous inflammatory cytokine signaling, JAK inhibition has been reported to attenuate chronic itch and improve skin barrier function by regulating the expression of skin barrier protein filaggrin.^{74, 75}

AIFA (Agenzia Italiana del Farmaco) and EMA (European Medicines Agency) recommendations for the use of all JAK inhibitors

AIFA has released the recommendations for the use of all JAK inhibitors for atopic dermatitis with the “determina” n. DG 63/2023 on 24 February 2023 («Aggiornamento scheda prescrizione cartacea dei farmaci JAKi (abrocitinib, upadacitinib) nel trattamento della dermatite atopica», available on Gazzetta Ufficiale della Repubblica italiana n. 58 del 9 marzo 2023) and the “Nota Informativa” of the 16 March 2023 (available at <https://www.aifa.gov.it/-/nota-informativa-importante-sugli-inibitori-della-janus-chinasi>). The same recommendations were previously released by EMA on 23 January 2023 (available at <https://www.ema.europa.eu/en/medicines/human/referrals/janus-kinase-inhibitors-jaki>) that endorsed the measures recommended by the Pharmacovigilance Risk Assessment Committee (PRAC) to minimize the risk of serious side effects (cardiovascular diseases, blood clots, cancer and serious infections) with Janus kinase (JAK) inhibitors used to treat several chronic inflammatory disorders. These risks were considered class effects and relevant with all approved indications of JAK inhibitors in inflammatory skin disorders. The AIFA’s recommendation were updated on 28 April 2023 [“Aggiornamento della scheda di prescrizione cartacea dei farmaci JAKi (abrocitinib, upadacitinib) nel trattamento della dermatite atopica. (Determina AIFA n. DG/197/2023) available on Gazzetta Ufficiale Serie Generale n.99 del 28-04-2023)]

According to the latest update Abrocitinib (50 mg and 100 mg) and upadacitinib (15 mg) are reimbursed for the treatment of severe atopic dermatitis (EASI score ≥ 24) in adult patients (the use for ≥ 12 years old adolescents is authorized but not reimbursed so far) who are candidates for systemic therapy:

- in the absence of EMA-indicated risk factors (elderly people over 65 years of age, subjects with increased risk of serious cardiovascular problems, such as myocardial infarction or stroke, long-term ex-smokers or smokers, and peoples at increased risk of cancer) if cyclosporine treatment has failed;
- in the presence of EMA- indicated risk factors: after failure, *i.e.* “inefficacy/loss of efficacy, occurrence of adverse events”, of reimbursed treatment options (ciclosporin and anti-interleukins) or the presence of factors that in the clinical judgement of the prescribing physician contraindicates or makes the above mentioned treatment options inappropriate in the individual patient”.

The AIFA’s recommendation also states that a lower dose, if available, may be recommended, depending on the medicinal product and the specific risk factor and that prescribers should inform patients of the risks associated with the use of JAK inhibitors.

On 7 July 2023, The AIFA’s Determina n. DG/267/2023 (Gazzetta Ufficiale Serie Generale n.157 del 07-07-2023) reported that oral baricitinib (2 mg and 4 mg) was reimbursed with the same clinical indications, limitations and recommendations reported above for abrocitinib and upadacitinib. In addition, it was added that baricitinib should be used in combination with topical corticosteroids.

JAK inhibitors can be prescribed only by dermatologists who work in centers specifically authorized by the Regional Health Authorities.

Abrocitinib

Abrocitinib is currently licensed for AE in those aged 12 and above in the United Kingdom. The EMA Committee for Medicinal Products for Human Use adopted a positive opinion on 14 October 2021, for adults only. As this approval came through after our consensus conferences, no consensus recommendation has been included in this iteration of the guideline.

Mechanisms of action and efficacy

Abrocitinib is an oral JAK1 selective inhibitor and has shown efficacy in patients with moderate-to-severe AE when used as a monotherapy (MONO-1 and -2 studies) and in combination with topical therapies in achieving treatment response in comparison with placebo (COMPARE study), as measured using IGA and EASI-75 response. For instance, the proportion of patients with EASI-75 response at week 12 was significantly higher with abrocitinib 100 mg (~40-45%) and abrocitinib 200 mg (~61-63%) compared with placebo (~10-12%) in the MONO studies. In the COMPARE study, the proportion of patients with EASI-75 response was significantly higher with abrocitinib 100 mg (~59%) and abrocitinib 200 mg (~70%) compared with placebo (27%).⁴ Similar efficacy has been demonstrated in the adolescent JADE TEEN trial for both the 100 mg and the 200 mg doses, in combination with topical therapies.⁷⁶ Importantly, in the COMPARE study (which had dupilumab as a comparator arm), higher responder rates were observed with abrocitinib 200 mg compared with dupilumab (*P*-values not calculated) after 16 weeks of treatment. The efficacy of abrocitinib 100 mg and dupilumab was similar in this subgroup. The results indicate that abrocitinib 200 mg may provide a higher probability of treatment response compared with dupilumab in patients with severe AE.⁷⁷

Dosage: acute flare, short term, long term

Abrocitinib is licensed at the 100 mg and the 200 mg daily doses, with the lower dose recommended for adolescents as a starting dose. One study assessed risk and probability of flares and recapture of treatment response following a flare. Of 1233 patients, 798 responders to induction with abrocitinib 200 mg (64.7%) were randomly assigned to dose maintenance, dose reduction or treatment withdrawal (placebo). The flare probability during maintenance was 18.9%, 42.6%, and 80.9% with abrocitinib 200 mg, abrocitinib 100 mg, and placebo, respectively, by week 52. Among patients with flare in

the abrocitinib 200 mg, abrocitinib 100 mg, and placebo groups, 36.6%, 58.8%, and 81.6% regained IGA 0/1 response, respectively, and 55.0%, 74.5%, and 91.8% regained EASI index response, respectively, with rescue treatment of abrocitinib 200 mg plus medicated topical therapy.⁷⁸

Safety

Based on long-term follow-up of patients from the phase II and III trials as well as one long-term extension study, with a total *n* of 2856 [1614 patient-years (PY)]; total exposure in the all-abrocitinib cohort was ≥ 24 weeks in 1248 patients and ≥ 48 weeks in 606 (maximum 108 weeks). In the placebo-controlled cohort (*N*=1540), dose-related adverse events (200 mg, 100 mg, placebo) were nausea (14.6%, 6.1%, 2.0%), headache (7.8%, 5.9%, 3.5%) and acne (4.7%, 1.6%, 0%). Platelet count was reduced transiently in a dose-dependent manner; two of 2718 patients (200-mg group) had confirmed platelet counts of $< 50 \times 10^3/\text{mm}^3$ at week 4. Incidence rates (IRs) were 2.33/100 PY and 2.65/100 PY for serious infection, 4.34/100PY and 2.04/100PY for herpes zoster, and 11.83/100 PY and 8.73/100 PY for herpes simplex in the 200-mg and 100-mg groups, respectively.⁷⁹

Monitoring

For baseline screening, the manufacturer's UK label laboratory monitoring recommendations are full blood count including platelet count, absolute lymphocyte count (ALC), absolute neutrophil count (ANC), and hemoglobin (Hb) as well as lipid parameters. A chest radiograph, creatinine phosphokinase level and an infection screening for HIV, hepatitis B and C as well as TB is advisable before initiation of therapy.

In practice, we recommend the same baseline screening and treatment monitoring investigations for all JAK inhibitors. For baseline screening, this is a full blood count, renal, liver and lipid profile as well as creatinine phosphokinase levels and hepatitis and TB screen, including a chest radiograph.

For monitoring purposes, we recommend a full blood count, renal, liver and lipid profile as well as creatinine phosphokinase level at 4 weeks into treatment and then three-monthly while on therapy.

Combination with other treatments

No studies assessing the use of abrocitinib with other systemic therapies have been published to date.

Special considerations

Abrocitinib is a new JAK inhibitor and has not been formally tested in other inflammatory diseases.


Cibinqo® (Pfizer Inc, New York, New York, U.S.A.) is indicated for the treatment of moderate to severe atopic dermatitis in adults who are candidates for systemic therapy. The recommended dose is 100 mg or 200 mg once daily based on individual patient presentation. A dose of 100 mg is recommended for patients at higher risk of venous thromboembolism (VTE), major adverse cardiovascular events (MACE) and malignancy. For patients ≥65 years of age, the recommended dose is 100 mg once daily. A dose of 200 mg once daily may be appropriate for patients with high disease burden who are not at higher risk of VTE, MACE and malignancy or patients with an inadequate response to 100 mg once daily. The lowest effective dose to maintain response should be used.

In patients with moderate to severe renal impairment a dose of 50 mg to 100 mg once daily is recommended. The drug is available in the following pharmaceutical forms: 50, 100, 200 mg tablets.

Reimbursement

Abrocitinib [Cibinqo® compresse (15 mg)] is reimbursed in Italy for the treatment of severe atopic dermatitis (EASI ≥24) in adult patients who are candidates for systemic therapy with the clinical indications, limitations and recommendations that are described in the box above (Determina AIFA n. DG/197/2023) available on Gazzetta Ufficiale Serie Generale n.99 del 28-04-2023).

Baricitinib

We recommend baricitinib in AE patients who are candidates for systemic treatment.	↑↑	<div style="text-align: center;">  <p>>75% (15/16)²</p> </div> Evidence and consensus based, see Evidence Report
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Baricitinib: in license for adults; dosage adults: 4 mg per day, reduction to 2 mg per day possible, depending on treatment response
 Certainty of evidence:^{4,5}
 Short term (8-16 weeks) vs. placebo (NMA main analysis)
 ⊕⊕⊕⊕ MODERATE for mean difference/standardized mean difference EASI, DLQI
 ⊕⊕⊕⊕ MODERATE - ⊕⊕⊕⊕ LOW for undesirable effects
 For baricitinib versus other drugs, see Evidence Report
¹One abstention.

Mechanisms of action and efficacy

Baricitinib is an oral selective JAK1 and JAK2 inhibitor. The drug has been tested in one phase 2 and several

phase 3 trials in adults with moderate-to- severe AE at 1, 2 and 4 mg once daily against placebo, showing significant improvement with regard to EASI from baseline to 16 weeks, in particular in the two higher doses {2 mg daily [mean difference, 5.6-point reduction; 95% CI, 0.4–10.9 (GRADE assessment: moderate certainty)] and 4 mg daily [mean difference, 5.2- point reduction; 95% CI, 0.1–10.4 (GRADE assessment: moderate certainty)]}.⁴ Similar efficacy has been shown in these studies with regard to the IGA and itch scores. The concomitant use of topical corticosteroids was allowed in one trial.⁸⁰

Dosage: acute flare, short term, long term

At present, Baricitinib data are available up to 52 weeks of follow-up,⁸¹ demonstrating sustained efficacy. There is no study that has looked at acute flare treatment and the paediatric study programme is still underway⁸² and no clear dosing guidance for paediatric patients is currently available.

Safety

The most common side-effects with baricitinib in clinical trials include an increase in LDL cholesterol, upper respiratory tract infections and headache. Acne is less common than with other JAK inhibitors. Infections reported with baricitinib include herpes simplex. However, the rate of these events reported in a recent combined safety study including 2531 patients from eight RCTs who were given baricitinib for 2247 patient-years (median duration 310 days) was overall low: eczema herpeticum (N.=11), cellulitis (N.=6) and pneumonia (N.=3). There were four opportunistic infections reported.⁸³ A transient increase in CPK may be seen, especially after extensive bodily exercise. No malignancies, gastrointestinal perforations, positively adjudicated cardiovascular events or tuberculosis were reported in the placebo-controlled period in baricitinib-treated patients. The frequency of herpes simplex was higher in the 4 mg group (6.1%) compared to the 2 mg (3.6%) and placebo groups (2.7%). Long-term safety data beyond 16 weeks are available from an integrated data base covering mostly rheumatoid arthritis patients for up to 9.3 years of treatment.⁸⁴

Monitoring

For baseline screening, the manufacturer advises that patients with suspected hepatitis B consult a liver specialist for advice before initiation of treatment. Lipid and liver

profiles need to be regularly monitored following treatment initiation. Screening for any hematological abnormalities is also advised.

In practice, we recommend the same baseline screening and treatment monitoring investigations for all JAK inhibitors. For baseline screening, this is a full blood count, renal, liver and lipid profile as well as creatinine phosphokinase levels and hepatitis and TB screen, including a chest radiograph.

For monitoring purposes, we recommend a full blood count, renal, liver and lipid profile as well as creatinine phosphokinase level at 4 weeks into treatment and then three-monthly while on therapy.

Combination with other treatments

No studies assessing the use of baricitinib with other systemic therapies in AE patients have been published to date, but the combination therapy with MTX is an established combination regimen in the management of rheumatoid arthritis.⁸⁵

Special considerations


AE patients with concomitant inflammatory diseases, such as rheumatoid arthritis, ankylosing spondylitis and psoriatic arthritis are likely to experience beneficial effects. Baricitinib is already licensed for this indication.

Olumiant® (Eli Lilly and Company, Indianapolis, Indiana, U.S.A.) is approved for the treatment of moderate to severe atopic dermatitis in adults who are candidates for systemic therapy. The recommended dose is 4 mg or 2 mg once daily based on individual patient presentation. A dose of 2 mg is recommended for patients at higher risk of venous thromboembolism (VTE), major adverse cardiovascular events (MACE) and malignancy. For patients ≥75 years of age, the recommended dose is 2 mg once daily. A dose of 4 mg once daily may be appropriate for patients with high disease burden who are not at higher risk of VTE, MACE and malignancy or patients with an inadequate response to 2 mg once daily. The lowest effective dose to maintain response should be used.

Reimbursement

Baricitinib [Olumiant® compresse 2 mg and 4 mg] is reimbursed in Italy (Gazzetta Ufficiale Serie Generale n.157 del 07-07-2023) with the same clinical indications, limitations and recommendations reported in the box above for all JAK inhibitors. Baricitinib should be used in combination with topical corticosteroids.

Upadacitinib

We recommend upadacitinib in AE patients who are candidates for systemic treatment.	↑↑	>50%  (8/15) Evidence and consensus based, see Evidence Report
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Upadacitinib: in license for ≥12 years; adults: 15 or 30 mg per day; age ≥65: 15 mg per day age 12-17 (≥30 mg/kg BW): 15 mg per day
 Certainty of evidence:^{4, 5}
 Short term (8-16 weeks) vs. placebo (NMA main analysis)
 ⊕⊕⊕⊕ LOW for mean difference **POEM**
 ⊕⊕⊕⊕ VERY LOW for undesirable effects
 For upadacitinib versus other drugs, see Evidence Report
 Upadacitinib is licensed for AE in adolescents (12 years and above) and adults.

Mechanisms of action and efficacy

Upadacitinib is a selective and reversible JAK inhibitor. There is one phase 2 trial including 167 adult patients that investigated three different doses of upadacitinib (30, 15 and 7.5 mg/day) for AE compared to placebo.⁸⁶ The trial was conducted over 16 weeks. Upadacitinib was superior to placebo for all dosage groups in EASI (mean change [SE] 74% [6.1%] for 30 mg, 62% [6.1%] for 15 mg, 39% [6.2%] for 7.5 mg and 23% [6.4%] for placebo [P=0.03, P<0.001, P<0.001]). There were also significant improvements seen with regard to the SCORAD index, NRS pruritus and POEM scores. The trials published since have shown similar efficacy.⁸⁷⁻⁸⁹

In a direct head-to-head trial enrolling adult AE patients randomized to receive upadacitinib (N.=348) and dupilumab (N.=344) 247 patients receiving upadacitinib (71.0%) and 210 patients receiving dupilumab (61.1%) achieved EASI-75 at 16 weeks (P=0.006). All ranked secondary end points also demonstrated the superiority of upadacitinib vs dupilumab, including improvement in Worst Pruritus NRS as early as week 1, achievement of EASI-75 as early as week 2, and EASI-100 at week 16. Rates of serious infection, eczema herpeticum, herpes zoster, and laboratory-related adverse events were higher for patients who received upadacitinib, whereas rates of conjunctivitis and injection-site reactions were higher for patients who received dupilumab.

Dosage: acute flare, short term, long term

Upadacitinib is licensed at the 15 and 30 mg doses for AE, and at 15 mg for rheumatoid arthritis, psoriatic arthritis and ankylosing spondylitis. Follow-up until week 52 is now available, showing long-term efficacy and safety profiles similar to the 16 week trials.⁹⁰ There is no study that has looked at acute flare treatment, and there are currently early phase AE trials in children >6 months.

Safety

The cumulative incidence rates of adverse events were 78.6% for 30 mg, 76.2% for 15 mg, 73.8% for 7.5 mg and 62.5% for placebo in the phase 2 trial and have been similar in the studies reported since.⁸⁶ Upper respiratory tract infections and acne were the most frequently reported adverse events for Upadacitinib. The cumulative incidence rates of severe adverse events were 0% for 30 mg, 2.4% for 15 mg, 4.8% for 7.5 mg and 2.4% for placebo. Low withdrawal rates were reported in the placebo and upadacitinib groups (N.<5 for each group).

Monitoring

The manufacturer advises that patients are screened for viral hepatitis B and C and TB. Lipid and liver profiles need to be measured at baseline and regularly following treatment initiation. Screening and monitoring for any haematological abnormalities are also advised, no later than 12 weeks.

In practice, we recommend the same baseline screening and treatment monitoring investigations for all JAK inhibitors. For baseline screening, this is a full blood count, renal, liver and lipid profile as well as creatinine phosphokinase levels and hepatitis and TB screen, including a chest radiograph.

For monitoring purposes, we recommend a full blood count, renal, liver and lipid profile as well as creatinine phosphokinase level at 4 weeks into treatment and then three-monthly while on therapy.

Combination with other treatments

No studies assessing the use of upadacitinib with other systemic therapies in AE patients have been published to date, but the combination therapy with MTX is an established combination regimen in the management of rheumatoid arthritis, albeit only with the 15 mg once a day dose.⁹¹

Special considerations

AE patients with concomitant inflammatory diseases, such as rheumatoid arthritis, psoriatic arthritis and ankylosing spondylitis are likely to experience beneficial effects, as upadacitinib is already licensed for this indication.⁹²

UPADACITINIB

Rinvoq® (Abbvie Inc, North Chicago, IL, USA) is indicated for the treatment of moderate to severe atopic dermatitis in adults and adolescents 12 years and older who are candidates for systemic therapy.

The recommended dose is 15 mg or 30 mg once daily based on individual patient’s presentation. A dose of 15 mg is recommended for patients at high risk of venous thromboembolism (VTE), major adverse cardiovascular events (MACE) and malignancy. A dose of 30 mg once daily may be appropriate for patients with high disease burden who are not at higher risk of VTE, MACE and malignancy or patients with an inadequate response to 15 mg once daily. The lowest effective dose to maintain response should be used. For adolescents (from 12 to 17 years of age) weighing at least 30 kg and for patients ≥65 years of age, the recommended dose is 15 mg once daily.


Upadacitinib should be used in the following patients only if no suitable treatment alternatives are available: those aged 65 years or above, those with history of atherosclerotic cardiovascular disease or other cardiovascular risk factors (such as current or past long-time smokers) and those with malignancy risk factors (such as current malignancy or history of malignancy).

Reimbursement

Upadacitinib (Rinvoq® compresse [15 and 30 mg]) is reimbursed in Italy for the treatment of severe atopic dermatitis (EASI ≥24) in adult patients who are candidates for systemic therapy with the clinical indications, limitations and recommendations that are described in the box above (Determina AIFA n. DG/197/2023) available on Gazzetta Ufficiale Serie Generale n.99 del 28-04-2023).

Other systemic treatment

Alitretinoin

<p>We suggest alitretinoin in AE patients with severe chronic hand eczema, who are candidates for systemic treatment, duly considering its teratogenicity.</p>	↑	<p style="text-align: center;">>75%</p>  <p style="text-align: center;">(14/15) Expert consensus</p>
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Alitretinoin: in label for adults with severe chronic hand eczema unresponsive to topical corticosteroids; dosage adults 10-30 mg per day

Mechanisms of action and efficacy

Alitretinoin is a retinoid binding both retinoic acid (RAR) and retinoic X (RXR) receptors, thus delivering anti-inflammatory and antiproliferative effects. It is licensed in some European countries for the treatment of chronic hand eczema irrespectively of its pathogenesis.

There is one large, multicenter randomized, placebo- controlled clinical trial involving 1032 patients with

chronic hand eczema, about one-third of which were probably atopic hand eczema patients.⁹³ Improvement of eczema was seen in 75% of the patients. The patient group suffering from atopic hand eczema was not analyzed separately, and extrapalmar symptoms have not been assessed in this trial.

Six patients with AE and prominent hand involvement were treated with alitretinoin for 12 weeks in an uncontrolled, open-label trial.⁹⁴ Both, palmar and extrapalmar lesions improved during the trial, as shown by the modified Total Lesion Symptom Score (mTLSS) hand eczema score and the SCORAD.

Dosage: acute flare, short term, long term

According to the mode of action, alitretinoin is suitable for long-term treatment. An alitretinoin treatment course should be planned for 3 to 6 months. The dosage of alitretinoin is 10–30 mg per day.

Safety

As alitretinoin is highly teratogenic, all women of child-bearing potential must adhere to a strict birth control program.

Monitoring

Before and during therapy: liver enzymes [aspartate aminotransferase (ASAT), alanine aminotransferase (ALAT), gamma-glutamyl transpeptidase (GGT)], cholesterol, triglycerides, basal thyroid stimulating hormone (TSH), free thyroxine (fT4) peripheral blood levels; pregnancy test in women with childbearing potential.

Combination with other treatments

Concomitantly to alitretinoin, topical therapy with corticosteroids, calcineurin inhibitors and emollients can be applied.

Special considerations

A retrospective analysis of children treated with alitretinoin because of hand eczema and other diagnoses including two severe AE patients, revealed that the response to alitretinoin was moderate in one subject, whereas the other patient failed to improve even after extending treatment to up to 11 months.⁹⁵

Alitretinoin is a class H drug in Italy and is approved for the treatment of chronic hand eczema, regardless of its atopic or non-atopic nature, resistant to potent topical cor-

ticosteroids. Patients must be enlisted in an AIFA registry monitoring the modalities of prevention of pregnancy and all female patients in child-bearing age must repeat a pregnancy test every month (https://www.aifa.gov.it/documenti/20142/1781182/Scheda_Registro_PPP_ind.Minjuvi_08.11.2022.zip)

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Conflicts of interest (authors of the Italian Adaptation of the EuroGuiDerm Guidelines)

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Authors' contributions

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