Policy Number	**SUR716.018**
Policy Effective Date	**09/01/2025**

Chemical Peels

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Related Policies (if applicable)
SUR717.001: Gender Assignment Surgery and Gender Reassignment Surgery with Related Services

Disclaimer

Carefully check state regulations and/or the member contract.
Each benefit plan, summary plan description or contract defines which services are covered, which services are excluded, and which services are subject to dollar caps or other limitations, conditions or exclusions. Members and their providers have the responsibility for consulting the member's benefit plan, summary plan description or contract to determine if there are any exclusions or other benefit limitations applicable to this service or supply. **If there is a discrepancy between a Medical Policy and a member's benefit plan, summary plan description or contract, the benefit plan, summary plan description or contract will govern.**

Coverage

Dermal chemical peels used to treat individuals with numerous (>10) actinic keratoses or other premalignant skin lesions, such that treatment of the individual lesions becomes impractical, **may be considered medically necessary**.

Epidermal chemical peels used to treat individuals with active acne that has failed a trial of topical and/or oral antibiotic acne therapy **may be considered medically necessary**. In this setting, superficial chemical peels with 40% to 70% alpha hydroxy acids are used as a comedolytic therapy. (Alpha-hydroxy acids can also be used in lower concentrations [8%] without the supervision of a physician.)

Epidermal chemical peels to treat photoaged skin, wrinkles, or acne scarring OR dermal chemical peels to treat end-state acne scarring **are considered cosmetic**.

Policy Guidelines

Requests for all chemical peels should be carefully evaluated to determine whether the treatment is primarily cosmetic. Epidermal peels would be considered medically necessary in individuals with active acne who have failed other therapy because active severe acne may lead to acne scarring and may be psychologically painful leading to low self-esteem, depression, and anxiety. Dermal peels would be considered medically necessary in individuals with multiple actinic keratoses because these premalignant lesions may warrant destruction or removal as an alternative to watchful waiting. Other applications of chemical peels, including treatment of photoaged skin, wrinkles, and acne scarring, are considered cosmetic.

Description

A chemical peel is a controlled removal of various layers of the skin with the use of a chemical agent. The most common use of chemical peeling is the treatment of photoaged skin. Chemical peeling has also been used for other conditions, including actinic keratoses, active acne, and acne scarring.

Chemical Peels

Chemical peels involve a controlled partial-thickness removal of the epidermis and the outer dermis. When skin is regenerated, a 2- to 3-mm band of dense, compact collagen is formed between the epidermis and the damaged layers of the dermis, resulting in the ablation of fine wrinkles and a reduction in pigmentation. These changes can be long-term, lasting 15 to 20 years and may be permanent in some individuals. Potential local complications include scarring, infection, hypopigmentation, hyperpigmentation, activation of herpes simplex, and toxic shock syndrome. (1)

Types of Peels

Chemical peels are often categorized by the depth of the peel: categories include superficial, medium-depth, and deep chemical peels. The precise depth of the peel depends on the concentration of the agent used, the duration of the application, and the number of applications. Possible indications for each type of peel and common chemicals used, as described by Cummings et al. (2005) (2) and others, is as follows.

Superficial Peels

Superficial peels (epidermal peels) affect the epidermis and the interface of the dermis-epidermis. This depth is considered appropriate for treating mild photoaging, melasma, comedonal acne, and postinflammatory erythema. Common chemical agents used for superficial peels include low concentrations of glycolic acid, 10% to 20% trichloroacetic acid (TCA), Jessner solution (a mixture of resorcinol, salicylic acid, lactic acid, and ethanol), tretinoin, and salicylic acid. As part of the treatment process, superficial peels generally cause mild erythema and desquamation, and healing time ranges from 1 to 4 days, depending on the strength of the chemical agent. With superficial peels, patients often undergo multiple sessions, generally, 6 to 8 peels performed weekly or biweekly.

Medium-Depth Peels
Medium-depth peels (dermal peels) extend into the epidermis to the papillary dermis. They are used for moderate photoaging, actinic keratoses, pigmentary dyschromias, and mild acne scarring. In the past, 50% TCA was a common chemical agent for medium-depth peels, but its use has decreased due to high rates of complications (e.g., pigmentary changes, scarring). Currently, the most frequently used agent is a combination of 35% TCA with Jessner solution or 70% glycolic acid. Phenol 88% alone is also used for medium-depth peels. The healing process involves mild-to-moderate edema, followed by the appearance of new, erythematous epithelium. Individuals are advised to wait at least 3 months before resuming skincare services (e.g., superficial chemical peels) and repeat medium-depth chemical peels should not be performed for at least 1 year.

Deep Peels
Deep chemical peels (another type of dermal peel) penetrate the mid-reticular dermis and have been used for patients with severe photodamage, premalignant skin neoplasms, acne scars, and dyschromias. The most common chemical agent used is Baker solution (which consists of 3 mL of 88% phenol, 8 drops of hexachlorophene [Septisol], 3 drops of croton oil, 2 mL of distilled water). The same depth can be achieved using 50% or greater TCA peel; however, the latter has a higher risk of scarring and pigmentation problems. Phenol is cardiotoxic, and patients must be screened for cardiac arrhythmias or medications that could potentially precipitate an arrhythmia. Phenol can also have renal and hepatic toxicities.

The likelihood and potential severity of adverse events increase as the strength of the chemicals and the depth of peels increases. With deep chemical peels, there is the potential for long-term pigmentary disturbances (i.e., areas of hypopigmentation), and selection of individuals willing to always wear makeup is advised. Moreover, chemical peels reduce melanin protection, so patients must use protective sunscreen for 9 to 12 months after a medium- to deep-facial peel.

<u>Applications</u>
Chemical peels are a potential treatment option for actinic keratoses and moderate-to-severe acne. Actinic keratoses are common skin lesions associated with extended exposure to the sun, with an estimated prevalence in the U.S. of 11% to 26%. (3) These lesions are generally considered to be a precursor of squamous cell carcinoma. (4) The risk of progression to invasive squamous cell carcinoma is unclear, but estimates vary from 0.1% to 20%. (3) For patients with multiple actinic keratoses, the risk of developing invasive squamous cell carcinoma is estimated as being between 0.15% and 80%. Treatment options include watchful waiting, medication treatment, cryosurgery, and surgical resection.

Acne vulgaris is the most common skin condition among adolescents, affecting an estimated 80% of teenagers aged 13 to 18 years old. (5) Acne, particularly moderate-to-severe manifestations, can cause psychologic distress including low self-esteem, depression, and anxiety. There are a variety of oral and topical treatments for acne.

Regulatory Status
U.S. Food and Drug Administration (FDA) clearance or approval of chemical agents used in peeling may not be relevant because these agents are prepared in-office, may have predated FDA approval, and/or may be considered cosmetic ingredients.

Rationale

Medical policies assess the clinical evidence to determine whether the use of technology improves the net health outcome. Broadly defined, health outcomes are the length of life, quality of life (QOL), and ability to function, including benefits and harms. Every clinical condition has specific outcomes that are important to patients and managing the course of that condition. Validated outcome measures are necessary to ascertain whether a condition improves or worsens; and whether the magnitude of that change is clinically significant. The net health outcome is a balance of benefits and harms.

To assess whether the evidence is sufficient to draw conclusions about the net health outcome of technology, 2 domains are examined: the relevance, and quality and credibility. To be relevant, studies must represent 1 or more intended clinical use of the technology in the intended population and compare an effective and appropriate alternative at a comparable intensity. For some conditions, the alternative will be supportive care or surveillance. The quality and credibility of the evidence depend on study design and conduct, minimizing bias and confounding that can generate incorrect findings. The randomized controlled trial (RCT) is preferred to assess efficacy; however, in some circumstances, nonrandomized studies may be adequate. RCTs are rarely large enough or long enough to capture less common adverse events and long-term effects. Other types of studies can be used for these purposes and to assess generalizability to broader clinical populations and settings of clinical practice.

Actinic Keratoses
Clinical Context and Therapy Purpose
The purpose of dermal chemical peels for individuals who have actinic keratosis is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations
The relevant population of interest is individuals with actinic keratosis.

Interventions
The therapy being considered is dermal chemical peels.

Comparators
The following therapies are currently being used to treat actinic keratosis: watchful waiting, medication treatment, cryosurgery, surgical resection, and photodynamic therapy.

Outcomes
The general outcomes of interest are destroying actinic keratosis, the durability of this effect, the development of cancerous lesions, QOL, and the harms of associated treatment-related morbidities.

The relevant follow-up is within weeks for the efficacy of treatment and years for the occurrence of cancerous lesions.

Study Selection Criteria
Methodologically credible studies for the indications within this review were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Systematic Reviews
Steeb et al. (2020) conducted a systematic review and meta-analysis assessing the efficacy and safety of chemical peels for the treatment of actinic keratosis. (6) A summary of the 8 trials included in the systematic review is shown in Table 1. This includes 4 RCTs, 2 non-randomized controlled trials, and 2 single-arm studies. Characteristics and results of the systematic review are summarized in Tables 2 and 3. Data analysis and interpretation of results were challenged by the presence of multiple study designs and the investigation of multiple distinct comparisons. The studies included in the review were at a high risk for selection bias because only one study clearly described the generation of a random sequence and performed allocation concealment. None of the patients in the studies were blinded; blinding of the outcome assessor was described in one study. Additionally, the chosen efficacy outcomes refer to short-term clearance rates but may not reflect long-term results. Overall, the authors concluded that additional high-quality studies and a standardization of peeling protocols were warranted in order to appropriately determine the value of chemical peeling as a treatment for actinic keratoses.

Table 1. Trials Included in a Systematic Review and Meta-Analysis of Chemical Peels for Actinic Keratosis

Trials	**Systematic Review**
	Steeb et al. (2020) (6)
Alfaro et al. (2012) (7)	●
Di Nuzzo et al. (2015) (8)	●
Holzer et al. (2017) (9)	●

Kaminaka et al. (2009) (10)	●
Lawrence et al. (1995) (11)	●
Marrero et al. (1998) (12)	●
Sandoval Osses et al. (2010) (13)	●
Sumita et al. (2018) (14)	●

Table 2. Summary of a Systematic Review and Meta-Analysis of Chemical Peels for Actinic Keratosis

Study	Dates	Trials	Participants	N (Range)	Design	Duration
Steeb et al. (2020) (6)	Until August 2019	8	Adults with a clinical or histopathological diagnosis of actinic keratosis	170 (13 to 32)	4 RCTs 2 non-randomized controlled trials 2 single-arm studies	NR

NR: not reported; RCT: randomized controlled trial.

Table 3. Results of a Systematic Review and Meta-Analysis of Chemical Peels for Actinic Keratosis

Study	Clearance Rate	Lesion-Specific Clearance	Mean Lesion Reduction Rate per Patient	Treatment-Related Pain (VAS)
Steeb et al. (2020) (6)				
TCA vs. PDT (n = 2 studies)				
Crude rate	0% (0/13) vs. 15.4% (2/13)[a]	66.1% (80/121) vs. 82.1% (101/123) 60.5% (214/354) vs. 82.6% (317/384)	65.9 ± 12.6 vs. 81.9 ± 12 51.1 ± 28.7 vs. 78.7 ± 26.2	7.31 ± 1.55 vs. 8.38 ± 1.56 5.1 ± 2.6 vs. 7.5 ± 2.3
Effect estimate	RR 0.20 (95% CI, 0.01 to 3.80)[a]	RR 0.75 (95% CI, 0.69 to 0.82)	MD -20.48 (95% CI, -31.55 to -9.41)	MD -1.71 (95% CI, -3.02 to -0.41)
TCA + Jessner's solution vs. 5-FU (n = 2 studies)				
Crude rate	15% (3/20) vs. 35% (7/20) 13.3% (2/15) vs. 46.7% (7/15)	81.7% (201/246) vs. 89% (202/227)	79.2 ± 19.5 vs. 89.6 ± 17.4	NR
Effect estimate	RR 0.36 (95% CI, 0.14 to 0.90)	RR 0.92 (95% CI, 0.85 to 0.99)[a]	MD -10.4 (95% CI, -23.63 to 2.83)[a]	NR
GA + 5-FU vs. GA (n = 1 study)				

Crude rate	22.2% (4/18) vs. 0% (0/18)	92.7% (217/234) vs. 15.8% (39/247)	92.1 ± 5.5 vs. 17.4 ± 8.7	NR
Effect estimate	RR 9.0 (95% CI, 0.52 to 155.86)	RR 5.87 (95% CI, 4.39 to 7.85)	MD 74.7 (95% CI, 69.95 to 79.45)	NR
Phenol peeling (n=1 study)				
Crude rate	90.62% (29/32)	NR	NR	NR
5-FU + GA (n = 1 study)				
Crude rate	30% (6/20)	92% (322/350)	NR	NR

[a] Only 1 study reported data for this outcome.
5-FU: 5-fluorouracil; CI: confidence interval; GA: glycolic acid; MD: mean difference; NR: not reported; PDT: photodynamic therapy; RR: risk ratio; TCA: trichloroacetic acid; VAS: visual analogue scale.

Section Summary: Actinic Keratoses

The evidence consists of a systematic review involving 8 studies - 4 RCTs, 2 non-randomized controlled trials, and 2 single-arm studies. Data analysis and interpretation of results were challenged by the high risk of bias of the primary studies, their imprecision, the variability of their peeling application protocols, and their focus on short-term clearance rates. Additional controlled studies, preferably randomized, are needed to determine the effect of chemical peels on the net health outcome in patients with actinic keratoses.

Moderate-to-Severe Active Acne

Clinical Context and Therapy Purpose

The purpose of epidermal chemical peels for individuals who have moderate-to-severe active acne is to provide a treatment option that is an alternative to or an improvement on existing therapies.

The following PICO was used to select literature to inform this policy.

Populations

The relevant population of interest is individuals with moderate-to-severe active acne.

Interventions

The therapy being considered is epidermal chemical peels.

Comparators

The following therapies are currently being used to treat active acne: topical or oral medications.

Outcomes

The general outcomes of interest are the resolution of severe acne and the harms of treatment-related morbidities.

The relevant follow-up is within weeks for the efficacy of treatment.

Study Selection Criteria

Methodologically credible studies for the indications within this review were selected using the following principles:

- To assess efficacy outcomes, comparative controlled prospective trials were sought, with a preference for RCTs.
- In the absence of such trials, comparative observational studies were sought, with a preference for prospective studies.
- To assess long-term outcomes and adverse events, single-arm studies that capture longer periods of follow-up and/or larger populations were sought.
- Studies with duplicative or overlapping populations were excluded.

Randomized Controlled Trials

RCTs comparing chemical peels to topical or oral medications for moderate-to-severe acne were not identified; the majority of studies evaluating the use of chemical peels for acne were in patients with mild-to-moderate disease. Of note, Kaminaka et al. (2014) conducted a double-blind, placebo-controlled randomized trial using a split-face design in Japan that evaluated 26 patients with moderate-to-severe facial acne. (15) Patients with moderate acne had 6 to 20 inflammatory lesions and up to 20 noninflammatory lesions; patients with severe acne had 21 to 50 inflammatory lesions. Failure of previous treatments was not an explicit inclusion criterion. Patients had to undergo a washout period of 2 months before study participation during which they could not use topical or oral antibiotics, retinoids, or corticosteroids. Participants then received a chemical peel treatment on a randomly selected side of the face, and a placebo peel on the other side of their face. Both treatments used the same pH acid gel vehicle (pH, 2.0) and the active treatment was a glycolic acid 40% peel. Treatments were given every 2 weeks for a total of 5 applications, and follow-up occurred 2 weeks after the last session (i.e., at 10-week follow-up). The overall therapeutic effect was judged by a blinded dermatologist as excellent or good for 23 (92%) of the chemical peel sides and 10 (40%) of the placebo sides; the difference between groups was statistically significant ($p<.01$). Moreover, there were statistically significant reductions in inflammatory lesions, and total lesion counts at each 2-week assessment and at the final 10-week assessment. No serious side effects or systemic adverse events were reported.

Section Summary: Moderate-to-Severe Active Acne

No RCTs comparing chemical peels to topical or oral medications in patients with moderate-to-severe acne were found. One placebo-controlled randomized trial was identified using a split-faced design with 26 patients who had moderate-to-severe acne. Outcomes (e.g., overall therapeutic effect) were significantly better in the chemical peel group. However, this trial testing a single chemical peel protocol in a relatively small number of patients provides insufficient evidence from which to draw conclusions about the safety and efficacy of chemical peels for treating active moderate-to-severe acne.

Summary of Evidence

For individuals who have actinic keratoses who receive dermal chemical peels, the evidence consists of a systematic review involving 8 studies - 4 randomized controlled trials (RCTs), 2 non-randomized controlled trials, and 2 single-arm studies. Relevant outcomes are symptoms, morbid events, quality of life, and treatment-related morbidity. Data analysis and interpretation of results were challenged by the high risk of bias of the primary studies, their imprecision, the variability of their peeling application protocols, and their focus on short-term clearance rates. Additional controlled studies, preferably randomized, are needed. The evidence is insufficient to determine that the technology results in an improvement in the net health outcome.

For individuals who have moderate-to-severe active acne who receive epidermal chemical peels, the evidence includes an RCT. Relevant outcomes are symptoms, morbid events, quality of life, and treatment-related morbidity. Results from the single, small, randomized, placebo-controlled, split-faced trial found greater efficacy with active treatment than with placebo. However, no studies were identified comparing chemical peel agents with conventional acne treatment. The evidence is insufficient to determine that the technology results in an improvement in the net health outcome.

Clinical Input

During a prior review, clinical input was consistently in agreement with the medically necessary indications for dermal and epidermal chemical peels.

Practice Guidelines and Position Statements

American Academy of Dermatology

In 2024, the American Academy of Dermatology (AAD) published guidelines on the management of acne vulgaris, which included the following statement on chemical peels (16): "Available evidence is insufficient to develop a recommendation on the use of...chemical peels (including glycolic acid, trichloroacetic acid, salicylic acid, Jessner's solution, or mandelic acid)...for the treatment of acne."

In 2021, the AAD published guidelines on the management of actinic keratosis, which gave a conditional recommendation based on moderate quality of evidence for the use of specific chemical peels for actinic keratosis. (17) The recommendation stated: "For patients with AKs [actinic keratosis], we conditionally recommend treatment with ALA [aminolevulinic acid]-red light PDT [photodynamic therapy] over trichloroacetic acid peel."

American Society for Dermatologic Surgery

In 2017, the American Society for Dermatologic Surgery published recommendations on the use of several skin treatments following a course of isotretinoin, a treatment for severe cystic acne. (18) Previously, a number of cosmetic skin treatments, including chemical peels, were discouraged for 6 months after the use of isotretinoin. These 2017 guidelines evaluated various treatments in the context of scarring and found that superficial chemical peels were safe as a treatment either concurrent with isotretinoin or within 6 months of its discontinuation. The lack of data on medium or deep chemical peels did not permit the Society to make a recommendation on those treatments.

Ongoing and Unpublished Clinical Trials

A currently ongoing/unpublished trial that might influence this policy is listed in Table 4.

Table 4. Summary of Key Trials

NCT Number	Trial Name	Planned Enrollment	Completion Date
NCT04429308	PDT Versus the Combination of Jessner's Solution and 35% TCA for Treatment of Actinic Keratoses on Upper Extremities: A Randomized Controlled Split-arm Trial	60	Dec 2025

NCT: national clinical trial.

Coding

Procedure codes on Medical Policy documents are included **only** as a general reference tool for each policy. **They may not be all-inclusive.**

The presence or absence of procedure, service, supply, or device codes in a Medical Policy document has no relevance for determination of benefit coverage for members or reimbursement for providers. **Only the written coverage position in a Medical Policy should be used for such determinations.**

Benefit coverage determinations based on written Medical Policy coverage positions must include review of the member's benefit contract or Summary Plan Description (SPD) for defined coverage vs. non-coverage, benefit exclusions, and benefit limitations such as dollar or duration caps.

CPT Codes	15788, 15789, 15792, 15793
HCPCS Codes	None

*Current Procedural Terminology (CPT®) ©2024 American Medical Association: Chicago, IL.

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Centers for Medicare and Medicaid Services (CMS)

The information contained in this section is for informational purposes only. HCSC makes no representation as to the accuracy of this information. It is not to be used for claims adjudication for HCSC Plans.

The Centers for Medicare and Medicaid Services (CMS) does not have a national Medicare coverage position. Coverage may be subject to local carrier discretion.

A national coverage position for Medicare may have been developed since this medical policy document was written. See Medicare's National Coverage at <https://www.cms.hhs.gov>.

Policy History/Revision	
Date	**Description of Change**
09/01/2025	Document updated with literature review. The following change was made to Coverage: Added "Epidermal chemical peels used to treat individuals with active acne that has failed a trial of topical and/or oral antibiotic acne therapy may be considered medically necessary. In this setting, superficial chemical peels with 40% to 70% alpha hydroxy acids are used as a comedolytic therapy. (Alpha-hydroxy acids can also be used in lower concentrations [8%] without the supervision of a physician.)" Added references 5, 15, 16, and 18.
10/15/2024	Document updated with literature review. Coverage unchanged. Reference 14 added.
01/01/2024	Reviewed. No changes.
07/01/2022	Document updated with literature review. Coverage unchanged. References 5-8 and 11-13 added.
07/01/2021	Reviewed. No changes.
01/15/2021	Document updated with literature review. Coverage unchanged. No new references added.
06/15/2019	Reviewed. No changes.
07/01/2018	Document updated with literature review. Coverage unchanged. References 3-5 were added; one reference was removed.
04/15/2017	Reviewed. No changes.
05/01/2016	Document updated with literature review. Coverage unchanged.
10/01/2015	Reviewed. No changes.
03/01/2014	Document updated with literature review. Coverage unchanged. CPT/HCPCS codes updated.
09/01/2009	Literature search, no change in coverage.
09/15/2007	Revised/Updated Entire Document
10/15/2008	Revised/Updated Entire Document
07/24/2004	Codes Revised/Added/Deleted
11/01/2000	Revised/updated entire document
09/01/1999	New medical document