

## Recommendations

---

### Standard guidelines of care for chemical peels

#### Niti Khunger

Member, IADVL Task Force\*, Department of Dermatology, Safdarjung Hospital, New Delhi, India

**Address for correspondence:** Dr. Niti Khunger, Department of Dermatology, Safdarjung Hospital, New Delhi, India.  
E-mail: [drniti@rediffmail.com](mailto:drniti@rediffmail.com)

---

#### ABSTRACT

Chemical peeling is the application of a chemical agent to the skin, which causes controlled destruction of a part of or the entire epidermis, with or without the dermis, leading to exfoliation and removal of superficial lesions, followed by regeneration of new epidermal and dermal tissues. **Indications** for chemical peeling include pigmentary disorders, superficial acne scars, ageing skin changes, and benign epidermal growths. **Contraindications** include patients with active bacterial, viral or fungal infection, tendency to keloid formation, facial dermatitis, taking photosensitizing medications and unrealistic expectations. **Physicians' qualifications:** The physician performing chemical peeling should have completed postgraduate training in dermatology. The training for chemical peeling may be acquired during post graduation or later at a center that provides education and training in cutaneous surgery or in focused workshops providing such training. The physician should have adequate knowledge of the different peeling agents used, the process of wound healing, the technique as well as the identification and management of complications. **Facility:** Chemical peeling can be performed safely in any clinic/outpatient day care dermatosurgical facility. **Preoperative counseling and informed consent:** A detailed consent form listing details about the procedure and possible complications should be signed by the patient. The consent form should specifically state the limitations of the procedure and should clearly mention if more procedures are needed for proper results. The patient should be provided with adequate opportunity to seek information through brochures, presentations, and personal discussions. The need for postoperative medical therapy should be emphasized. **Superficial peels** are considered safe in Indian patients. **Medium depth peels** should be performed with great caution, especially in dark skinned patients. **Deep peels** are not recommended for Indian skin. It is essential to do **prepeel priming** of the patient's skin with sunscreens, hydroquinone and tretinoin for 2-4 weeks. **Endpoints in peels:** For glycolic acid peels: The peel is neutralized after a predetermined duration of time (usually three minutes). However, if erythema or epidermolysis occurs, seen as grayish white appearance of the epidermis or as small blisters, the peel must be immediately neutralized with 10-15% sodium bicarbonate solution, regardless of the duration of application of the peel. The end-point is frosting for TCA peels, which are neutralized either with a neutralizing agent or cold water, starting from the eyelids and then the entire face. For salicylic acid peels, the end point is the pseudofrost formed when the salicylic acid crystallizes. Generally, 1-3 coats are applied to get an even frost; it is then washed with water after 3-5 minutes, after the burning has subsided. Jessner's solution is applied in 1-3 coats until even frosting is achieved or erythema is seen. Postoperative care includes sunscreens and moisturizers. Peels may be repeated weekly, fortnightly or monthly, depending on the type and depth of the peel.

**Key Words:** Glycolic acid, Trichloroacetic acid, Salicylic acid

\*The Indian Association of Dermatologists, Venereologists and Leprologists (IADVL) **Dermatosurgery Task Force** consisted of the following members: Dr. Venkataram Mysore (co-ordinator), Dr. Satish Savant, Dr. Niti Khunger, Dr. Narendra Patwardhan, Dr. Davinder Prasad, Dr. Rajesh Buddhadev, Lt. Col. Dr. Manas Chatterjee, Dr. Somesh Gupta, Dr. MK Shetty, Dr. Krupashankar DS, Dr. KHS Rao, Dr. Maya Vedamurthy, Ex offi-cio members: Dr. Chetan Oberai, President IADVL (2007-2008), Dr. Koushik Lahiri, Secretary IADVL, Dr. Sachidanand S, President IADVL (2008-2009), and Dr. Suresh Joshipura, Immediate Past president IADVL (2007-2008).

Evidence - Level A- Strong research-based evidence- Multiple relevant, high-quality scientific studies with homogeneous results, Level B- Moderate research-based evidence- At least one relevant, high-quality study or multiple adequate studies, Level C- Limited research-based evidence- At least one adequate scientific study, Level D- No research-based evidence- Based on expert panel evaluation of other information

For Disclaimers and Disclosures, please refer to the table of contents page (page 1) of this supplement.

The printing of this document was funded by the IADVL.

**How to cite this article:** Khunger N. Standard guidelines of care for chemical peels. Indian J Dermatol Venereol Leprol 2008;74:S5-S12.

**Received:** August, 2007. **Accepted:** May 2008. **Source of Support:** Nil. **Conflict of Interest:** Nil

Khunger N: Standard guidelines of care for chemical peels

## INTRODUCTION

The concept of peeling the skin to improve the texture, smoothen and beautify it has been used since ancient times. In ancient Egypt, Cleopatra used sour milk, now known to contain lactic acid, an alpha hydroxy acid while French women used old wine containing tartaric acid, to enhance the appearance of the skin.<sup>[1, 2]</sup> Chemical peeling is a common office procedure that has evolved over the years, using the scientific knowledge of wound healing after controlled chemical skin injury.<sup>[3]</sup> In spite of the advent of newer techniques and lasers, peeling has stood the test of time as a simple procedure, requiring hardly any instrumentation to rejuvenate the skin.

### Definition

Chemical peeling is the application of a chemical agent to the skin, which causes controlled destruction of a part or entire epidermis, with or without the dermis, leading to exfoliation, removal of superficial lesions, followed by regeneration of new epidermal and dermal tissues.

## RATIONALE AND SCOPE

These guidelines identify the indications for chemical peels, various agents that can be utilized, methodology, pre- and postpeel care, associated complications, and expected results.

## INDICATIONS OF CHEMICAL PEELS<sup>[1,4]</sup>

### *Pigmentary disorders*

- Melasma
- Postinflammatory hyperpigmentation
- Freckles
- Lentigines
- Facial melanoses

### *Acne*

- Superficial acne scars
- Postacne pigmentation
- Comedonal acne
- Acne excoriée
- Acne vulgaris-mild to moderately severe acne

### *Aesthetic*

- Photoaging
- Fine superficial wrinkling
- Dilated pores
- Superficial scars

### *Epidermal growths*

- Seborrheic keratoses
- Actinic keratoses
- Warts
- Milia
- Sebaceous hyperplasia
- Dermatoses papulosa nigra

## CONTRAINDICATIONS<sup>[1,4]</sup>

- i. Active bacterial, viral, fungal or herpetic infection
- ii. Open wounds
- iii. H/O (history of) drugs with photosensitizing potential
- iv. Preexisting inflammatory dermatoses such as psoriasis, atopic dermatitis
- v. Uncooperative patient (patient is careless about sun exposure or application of medicine)
- vi. Patient with unrealistic expectations.
- vii. For medium depth and deep peels-history of abnormal scarring, keloids, atrophic skin, and isotretinoin use in the last six months.<sup>[5]</sup>

## PHYSICIANS' QUALIFICATIONS

### 1. General

- a. The physician should be a trained dermatologist.
- b. The physician should have knowledge of the skin and subcutaneous tissue, including structural and functional differences and variations in skin anatomy of the facial cosmetic unit.

### 2. Specific

- a. The physician should have appropriate training in chemical peeling either during postgraduation or later at a center that routinely provides education and training in cutaneous surgery. Such training may also be obtained in focused workshops providing such training.
- b. The physician should have knowledge of the basic chemistry of peels, such as acids, bases, pH and pK<sub>a</sub> of peeling solutions and the mechanism of action of peels.<sup>[5]</sup> Familiarity with the properties of each peeling agent to be used is critical for successful outcome.
- c. The physician should know all aspects of mechanism of wound healing after chemical skin injury.
- d. The physician should be well versed with all aspects of pathogenesis and the medical therapy of the condition to be peeled, such as melasma, acne,

- photodamage etc.
- e. The physician should be well versed with early recognition, prevention and treatment of postoperative complications such as prolonged erythema, postinflammatory hyperpigmentation, impending scarring etc.

## PREPEEL ASSESSMENT

- A. History should include general medical history, degree of sun exposure, occupation to judge the level of sun exposure, history of herpes simplex, recent isotretinoin treatment in the last six months (for medium depth and deep peels), keloidal tendency, tendency for postinflammatory hyperpigmentation, current medications, any previous surgical treatment, immunocompromising conditions, and smoking (may delay healing in deep peels; this is not relevant for superficial peels). In patients in whom phenol peels are planned, history of systemic disease, particularly cardiac disease, should be taken.
- B. Detailed medical examination should include general physical and cutaneous examination including skin type, degree of photoaging, degree of sebaceous activity (oily or dry skin), presence of postinflammatory hyperpigmentation, keloid or hypertrophic scar, infection, and preexisting inflammation.
- C. Investigations
- Skin biopsy** should be done when indicated, to confirm diagnoses and see the level of pigmentation, *e.g.*, mixed or dermal melasma, lichen planus pigmentosus. No specific investigations are indicated for superficial peels.
  - In patients in whom deep (phenol) peels are planned, hemogram, urinalysis, liver and renal function tests and electrocardiograph may be carried out as cardiac complications such as life-threatening arrhythmia, are recognized as complications of deep peeling.
- D. Documentation
- Informed consent after counseling as below
  - Photographic record
- Counseling:** Proper counseling is very important and should include:
- Evaluation of the psychological aspects to judge the motivation and expectations of the patient.
  - Explanation that patient should have realistic expectations; this is particularly important in the media-hyped patient who may have unrealistic expectations.

- Explanation about the nature of treatment, expected outcome. It is advisable to downplay the degree of improvement expected.
- Information about the time taken for recovery of normal skin and importance of maintenance regimens.
- Discussion of side effects, likely and unlikely complications, and particularly, pigmentation changes.

## Preprocedure treatment recommendations (Priming):<sup>[1,5,6]</sup>

Priming is essential for at least 2-4 weeks prior to the procedure. Priming helps to reduce wound healing time, facilitates uniform penetration of peeling agent, detects intolerance to any agent, enforces patient compliance and reduces the risk of complications.

- Control any active infection or preexisting dermatoses.
- Broad-spectrum sunscreens.
- Hydroquinone (2-4%) in patients prone to postinflammatory hyperpigmentation.
  - Patients may also be primed at home by using mild topical peeling agents such as tretinoin 0.025%, adapalene 0.1%, Glycolic acid 6-12%, kojic acid, azelaic acid, etc (agents which are likely to be used in postprocedure maintenance). Tretinoin is known to reduce healing time after resurfacing.<sup>[7]</sup> The choice of the priming agent depends on the individual physician's preference and individualized patient requirements.
  - In patients with history of herpes simplex posted for medium depth and deep peels, antiviral therapy with acyclovir or famciclovir is recommended, beginning two days prior to the procedure and continued for 7-10 days until complete reepithelialization.

## REAGENTS

- Correctly labeled peeling agents in various concentrations
- Alcohol to clean the skin
- Acetone to degrease the skin
- Cold water
- Syringes filled with normal saline for irrigation of the eyes, in case of accidental spillage.
- Neutralizing solutions: Specific neutralizers are mentioned under "description of individual peels."

Khunger N: Standard guidelines of care for chemical peels

## EQUIPMENT

- i. Glass cup or beaker in which the required agent is poured
- ii. Head band or cap for the patient
- iii. Gloves
- iv. Cotton-tipped applicators or swab sticks
- v. 2" x 2" cotton gauze pieces
- vi. Fan for cooling
- vii. Timer for alpha-hydroxy acid peels

## PEELING AGENTS

1. Alpha-hydroxy acids, AHA Monocarboxylic acids: Glycolic acid (Level A)<sup>[8-14]</sup>, Lactic acid<sup>[15]</sup> (Level B), Bicarboxylic acid: Malic acid (Level C), Tricarboxylic acid: Citric acid (Level C)
2. Beta-hydroxy acids, BHA (salicylic acid)<sup>[16-19]</sup> (Level A)
3. Trichloroacetic acid (TCA)<sup>[20-22]</sup> (Level A)
4. Alpha-keto acids (pyruvic acid)<sup>[23]</sup> (Level C)
5. Resorcinol (Level B)
6. Jessner's solution<sup>[24, 25]</sup> (Salicylic acid 14 g, Lactic acid 14 g, Resorcinol 14 g with Ethanol to make 100 mL) (Level B)
7. Retinoic acid<sup>[26]</sup> (Level C)
8. Phenol<sup>[27-30]</sup> Type I-II skin (Level A) Type III-IV skin (Level C)

Classification of peels according to the histological depth of necrosis:<sup>[5]</sup>

- A. **Very Superficial light peels:** Necrosis up to the level of stratum corneum. Agents used: TCA 10%, GA 30-50%, Salicylic acid 20-30%, Jessner's solution 1-3 coats, Tretinoin 1-5%
- B. **Superficial light peels:** Necrosis through the entire epidermis up to basal layer. Agents used: TCA 10-30%, GA 50-70%, Jessner's solution 4-7 coats
- C. **Medium depth peels:** Necrosis up to upper reticular dermis. Agents used: TCA 35-50%, GA 70% plus TCA 35%, 88% phenol un-occluded, Jessner's solution plus TCA 35%, solid CO<sub>2</sub> plus TCA 35%
- D. **Deep peels:** Necrosis up to mid-reticular dermis. Agents used: Baker-Gordon phenol peel

## RECOMMENDATIONS

Anesthesia: Anesthesia is not required in superficial and medium depth peels. Mild tranquilizers or anxiolytics may be used in anxious patients.

**Safety Precautions before peeling:** The label on the bottle must be checked before applying the peel; the head should be elevated to 45°. To avoid accidental spillage, the open bottle or the soaked applicator should not be passed over the face. A syringe filled with water or saline should be kept ready for irrigation of the eyes in case of accidental spillage.

**Skin preparation before peeling**

- i. The patient is asked to wash the face with soap and water.
- ii. The hair is pulled back with a hair band or cap.
- iii. The patient lies down with head elevated to 45° with the eyes closed.
- iv. Using 2" x 2" gauze pieces, the skin is cleaned with alcohol and then degreased with acetone.

## PROCEDURE FOR SUPERFICIAL PEELS<sup>[1,4,8]</sup>

1. The required strength of the peeling agent is poured into a glass beaker and the neutralizing agent is also kept ready.
2. Sensitive areas like the inner canthus of the eyes and nasolabial folds are protected with Vaseline.
3. The peeling agent is then applied either with a brush or cotton-tipped applicator or gauze.
4. The chemical is applied quickly as cosmetic units on the entire face, beginning from the forehead, then the right cheek, nose, left cheek and chin in that order. If required, the perioral, upper and lower eyelids are treated last. Feathering strokes are applied at the edges to blend with surrounding skin and prevent demarcation lines.
5. For glycolic acid peels, the peel is neutralized after the predetermined duration of time (usually three minutes). However, if erythema or epidermolysis occurs, seen as grayish white appearance of the epidermis or small blisters, the peel must be neutralized immediately irrespective of the duration. Neutralization is done with 10-15% sodium bicarbonate solution or neutralizing lotion and then, washed off with water.
6. For TCA peels, the end-point is frosting and neutralization is either with a neutralizing agent or cold water, starting from the eyelids and then the entire face.
7. When the salicylic acid peel is applied, it crystallizes forming a pseudo-frost; generally, 1-3 coats are applied to get an even frost. It is then washed with water after 3-5 minutes, after the burning subsides.
8. Jessner's solution is applied in 1-3 coats to get even

- frosting; the endpoint is erythema or even frosting.
9. A cooling fan helps to reduce burning of the skin.
  10. The skin is gently dried with gauze and the patient is asked to wash with cold water until the burning subsides. The face is patted dry; rubbing should be avoided.
  11. Tretinoin peels are yellow peels that are left on for 4-5 hours and then washed away.
  12. Very superficial peels may be repeated every 1-2 weeks and superficial peels every 2-4 weeks.

## MEDIUM DEPTH AND DEEP PEELS

Medium depth peels should be done with great caution in dark skinned patients because of the high risk of prolonged hyperpigmentation.<sup>[1, 31]</sup> Deep phenol peels are not recommended for dark skins of types IV-VI because of high risk of prolonged or permanent pigmentary changes,<sup>[1]</sup> although modified phenol peels are being used in types III-IV Asian skins.<sup>[28-30]</sup>

## POSTOPERATIVE CARE

The aim of good postoperative care is to prevent or minimize complications and ensure early recovery. This is most important in dark skinned patients in whom pigmentary alterations are common. A careful maintenance program is essential to maintain the results of chemical peeling in most patients.

- i. In the postpeel period, edema, erythema and desquamation occur. In superficial peels, this lasts for 1-3 days, whereas in deeper peels, it lasts for 5-10 days.
- ii. Mild soap or a non-soap cleanser may be used. If there is crusting, a topical antibacterial ointment should be used to prevent bacterial infection.
- iii. Clear instructions must be given to the patient for the postprocedure period.
- iv. Cold compresses or calamine lotion may be used to soothe the skin.
- v. They should be told to use broad-spectrum sunscreens and only bland moisturizers until peeling is complete.
- vi. They should avoid peeling or scratching the skin.
- vii. Analgesics are not usually needed but may be advised in case of burning sensation.

## COMPLICATIONS<sup>[1,32-35]</sup>

Proper patient selection, adequate counseling, priming the

skin and supportive medical therapy in addition to good intra- and postoperative care, are essential for satisfactory cosmetic results. The best way to avoid complications is to identify patients at risk and use lighter peels. The deeper the peel, the greater is the risk of complications. The patients at risk are those with a history of postinflammatory hyperpigmentation, keloid formation, heavy occupational exposure to sun such as field workers, uncooperative patients and patients with a history of sensitive skin who are unable to tolerate sunscreens, hydroquinone etc.

- i. Pigmentary changes: Postinflammatory hyperpigmentation and hypopigmentation. These can be very persistent and often difficult to treat. They may be treated with broad-spectrum sunscreens, topical corticosteroids, tretinoin, hydroquinone or alpha-hydroxy acids.
- ii. Infection: Bacterial (*Staphylococcus*, *Streptococcus*, *Pseudomonas*), viral (Herpes simplex) and fungal (Candida). They should be treated aggressively and appropriately.
- iii. Scarring is rare in superficial peels. Proper priming, proper choice of peeling agent and postoperative care can help in prevention of this complication.
- iv. Allergic reactions
- v. Milia
- vi. Acneiform eruptions
- vii. Lines of demarcation
- viii. Textural changes
- ix. Persistent erythema: Erythema persisting for more than three weeks after a peel, is indicative of early scarring and should be treated with potent topical corticosteroids for  $\leq 2$  weeks.
- x. Toxicity: Although rare, it may occur with resorcinol, salicylic acid and phenol.<sup>[5]</sup>

## COMBINATION PEELS AND PROCEDURES

- A. combination of peeling agents enhances the depth of the peel without using a higher concentration of the peeling agent. However, these medium depth peels should be used cautiously in darker skinned patients because of the risk of uneven depth of peeling and increased risk of side effects, such as postinflammatory hyperpigmentation and scarring.
  - i. Glycolic acid 70% combined with TCA 35% (Coleman's Peel)<sup>[36]</sup> (level C). In darker skins, lower concentrations of TCA (10-25%) may be used (level D).<sup>[1]</sup>
  - ii. Solid CO<sub>2</sub> combined with 35% TCA (Brody's peel) (level C).

Khunger N: Standard guidelines of care for chemical peels

- iii. Jessner's solution with 35% TCA (Monheit's Peel) (level C).<sup>[37]</sup>
- B. Two procedures can also be combined to blend cosmetic units and avoid demarcation lines:<sup>[38-40]</sup>
  - i. Chemical peeling combined with dermabrasion: This procedure was originally used by combining application of 50% TCA followed by dermabrasion for post-acne scarring. However, 50% TCA causes scarring and its use is not advocated anymore.
  - ii. Chemical peeling can also be combined with

laser resurfacing for skin rejuvenation. First, a chemical peel is performed and then, the deeper wrinkles in the periorbital and perioral areas are treated with pulsed CO<sub>2</sub> laser (level C).

- iii. Chemical Peel with dermasanding using sandpaper (level C).
- iv. Chemical peeling with Botulinum Toxin (level C).
- v. Chemical peeling with fillers (level C).

## CONCLUSIONS

Chemical peeling is a simple office procedure used for the treatment of dyschromias, photoaging, and superficial scarring that can lead to excellent cosmetic improvement, when repeatedly performed in carefully selected patients. Although various depths of peels have been described, superficial and medium depth peels are safer for Indian patients. Deep chemical peels should be avoided because of the risk of permanent pigmentary changes. The type, depth and concentration of the peel should be selected according to the pathology of the condition (Table 1). Chemical peels are not one-time procedures and should be repeated with maintenance peels to achieve maximum improvement and prevent recurrence. With the advent of lasers and newer techniques, the use of chemical peels has

**Table 1A: A useful classification of peels, peeling agents and indications in Indian skin**

Peel depth	Level of peel	Peeling agent	Indications
Very superficial (exfoliation)	Up to stratum corneum	GA 30-50% TCA 10% Jessner's Solution (1-3 coats) Salicylic acid (20-30%)	Active acne
Superficial (epidermal)	Up to basal layer	GA 50-70% TCA 10-30% Jessner's Solution (4-7 coats)	Ephelides Epidermal melasma
Medium (papillary dermal)	Up to papillary dermis	GA 70% TCA 35-50%	Lentiginos Dermal melasma

**Table 1B: Comparison between commonly used peeling agents**

Agent	Advantages	Disadvantages
Glycolic acid	<ul style="list-style-type: none"> <li>• Even very superficial peels achieve significant results</li> <li>• Safe and effective at low concentrations</li> <li>• No systemic toxicity</li> <li>• Long shelf life</li> </ul>	<ul style="list-style-type: none"> <li>• Results not always predictable. Great variability in reactivity and efficacy</li> <li>• Tendency to penetrate unevenly.</li> <li>• Endpoint difficult to judge, greater chances of overpeeling.</li> <li>• Dermal wounds and scarring can occur.</li> <li>• Has to be neutralized.</li> <li>• Difficult to prepare and obtain standardized solutions.</li> <li>• Expensive</li> </ul>
Trichloroacetic acid	<ul style="list-style-type: none"> <li>• Peel depth correlates with intensity of skin frost.</li> <li>• Easy to visualize and apply evenly.</li> <li>• Endpoint easy to judge.</li> <li>• No need to neutralize.</li> <li>• No systemic toxicity.</li> <li>• Inexpensive</li> <li>• Easy to procure and prepare</li> </ul>	<ul style="list-style-type: none"> <li>• Pigmentary changes are common</li> <li>• Scarring can occur with high concentration.</li> <li>• Limited shelf life.</li> </ul>
Salicylic acid	<ul style="list-style-type: none"> <li>• Superficial peeling agent with a predictable response.</li> <li>• Safe in all skin types I-VI.</li> <li>• Lipophilic, hence very effective for acne, oily skin.</li> <li>• Causes a white pseudofrost, hence, easy to visualize and apply evenly.</li> <li>• Endpoint easy to judge.</li> <li>• No need to neutralize.</li> <li>• Does not penetrate deeply.</li> <li>• Safer, low incidence of significant complications.</li> <li>• Has anesthetic effect.</li> <li>• Easy to procure and prepare.</li> <li>• Inexpensive</li> </ul>	<ul style="list-style-type: none"> <li>• Can be absorbed systemically, when applied over large areas in high concentrations, causing salicylism.</li> <li>• Contraindicated in patients allergic to aspirin and during pregnancy and lactation.</li> <li>• Limited depth of peeling.</li> <li>• Minimal efficacy in severe photodamaged skin.</li> </ul>

declined; however, its simplicity as an office procedure, minimal morbidity, easy availability and cost-effectiveness ensure that it still holds an important place as a tool to treat dyschromias and photoaging. Careful patient selection, priming of the skin, standardization of peels, postpeel care and maintenance programs are essential to achieve excellent cosmetic results.

## REFERENCES

1. Savant SS. Superficial and medium depth chemical peeling. In: Text book of dermatosurgery and cosmetology. In: Savant SS, editor. 2<sup>nd</sup> ed. ASCAD; 2005. p. 177-95
2. Brody H. History of chemical peels. In: Baxter S, editor. Chemical peeling and resurfacing. 2nd ed. St. Louis: Mosby Year Book Inc; 1997. p. 1-5.
3. Stegman SJ. A comparative histologic study of the effects of the three peeling agents and dermabrasion on normal and sun damaged skin. *Aesth Plast Surg* 1982;6:123-35.
4. Monheit GD, Kayal JD. Chemical peeling. In *Techniques of Dermatologic Surgery*. Nouri K, Leal-Khoury, editors. Elsevier; 2003. p. 233-44.
5. Chemical Peels. In: Rubin MG, editor. *Procedures in cosmetic dermatology*. Elsevier Inc.; 2006 p. 1-12.
6. Roenigk RK, Retinoids, dermabrasion, chemical peel and keloids. In: Roenigk RK, Roenigk HH Jr, editors. *Surgical Dermatology: advances in current practice*. St Louis: Mosby; 1993. p. 376-83.
7. Hevia O, Nemeth AJ, Taylor JR. Tretinoin accelerates healing after TCA chemical peels. *Arch Dermatol* 1991;127:678-82.
8. Rubin MG. Glycolic acid peels. *Manual of chemical peels-superficial and medium depth*. In: Rubin MG, editor, 1st ed. Philadelphia: JB Lippincot Co; 1995. p. 89-102.
9. Ditre CM. Alpha-hydroxy acid peels. *Chemical Peels. Procedures in cosmetic dermatology*. In: Rubin MG, editor. Elsevier Inc.; 2006. p. 27-40.
10. Grover C, Reddy BS. The therapeutic value of glycolic acid peels in dermatology. *Indian J Dermatol Venereol Leprol* 2003;69:148-50.
11. Javaheri SM, Handa S, Kaur I, Kumar B. Safety and efficacy of glycolic acid facial peel in Indian women with melasma. *Int J Dermatol* 2001;40:354-7.
12. Wang CM, Huang CL, Hu CT, Chan HL. The effect of glycolic acid on treatment of acne in Asian skin. *Dermatol Surg* 1997;23:23-9.
13. Lim JTE, Tham SN. Glycolic acid peels in the treatment of melasma among Asian women. *Dermatol Surg* 1997;23:177-9.
14. Burns RL, Prevost-Blank PL, Lawry MA, Lawry TB, Faria DT, Fivenson DP. Glycolic acid for post inflammatory hyperpigmentation in black patients. *Dermatol Surg* 1997;23:171-5.
15. Sharquie KE, Al-Tikreety MM, Al-Mashhadani SA. Lactic acid as a new therapeutic peeling agent in melasma. *Dermatol Surg* 2005;31:149-54.
16. Vedamurthy M. Salicylic acid peels. *Indian J Dermatol Venereol Leprol* 2004;70:136-8.
17. Lee HS, Kim IH. Salicylic acid peels for the treatment of acne vulgaris in Asian patients. *Dermatol Surg* 2003;5:1196-9.
18. Grimes PE. Salicylic acid peels. *Chemical Peels. Procedures in cosmetic dermatology*. In: Rubin MG, editor. Elsevier Inc.; 2006. p. 47-55.
19. Bari AU, Iqbal Z, Rahman SB. Tolerance and safety of superficial chemical peeling with salicylic acid in various facial dermatoses. *Indian J Dermatol Venereol Leprol* 2005;71:87-90.
20. Leonhardt JM, Lawrence N. Trichloroacetic acid (TCA) peels. *Chemical Peels. Procedures in cosmetic dermatology*. In: Rubin MG, editor. Elsevier Inc.; 2006. p. 73-86.
21. Chun EY, Lee JB, Lee KH. Focal trichloroacetic acid peel method for benign pigmented lesions in dark-skinned patients. *Dermatol Surg* 2004;30:512-6.
22. Rubin MG. Trichloroacetic acid peels. *Manual of chemical peels-superficial and medium depth*. In: Rubin MG, editor, 1st ed. Philadelphia: JB Lippincot Co.; 1995. p. 110-29.
23. Ghersetich I, Brazzini B, Peris K, Cotellessa C, Manunta T, Lotti T. Pyruvic acid peels for the treatment of photoaging. *Dermatol Surg* 2004;30:32-6.
24. Rubin MG. Jessner's peels. *Manual of chemical peels-superficial and medium depth*. In: Rubin MG, editor, 1<sup>st</sup> ed. Philadelphia: JB Lippincot Co.; 1995. p. 79-88.
25. Fulton JE Jr. Jessner's Peel. *Chemical Peels. Procedures in cosmetic dermatology*. In: Rubin MG, editor, Elsevier Inc. 2006. p. 57-71.
26. Khunger N, Sarkar R, Jain RK. Tretinoin peels versus glycolic acid peels in the treatment of Melasma in dark-skinned patients. *Dermatol Surg* 2004;30:756-60.
27. Stone PA. Phenol peeling. *Chemical Peels. Procedures in cosmetic dermatology*. In: Rubin MG, editor, Elsevier Inc.; 2006. p. 87-114.
28. Piamphongsant T. Phenol-castor oil: Modified peel for dermal melasma. *Dermatol Surg* 2006;32:611-7.
29. Park JH, Choi YD, Kim SW, Kim YC, Park SW. Effectiveness of modified phenol peel (Exoderm) on facial wrinkles, acne scars and other skin problems of Asian patients. *J Dermatol* 2007;34:17-24.
30. Landau M. Cardiac complications in deep chemical peels. *Dermatol Surg* 2007;33:190-3.
31. Al-Waiz MM, Al-Sharqi AI. Medium-depth chemical peels in the treatment of acne scars in dark-skinned individuals. *Dermatol Surg* 2002;28:383-7.
32. Resnick SS, Resnick BI. Complications of chemical peeling. *Dermatol Clin* 1995;13:309-12.
33. Duffy DM. Avoiding complications. *Chemical Peels. Procedures in cosmetic dermatology*. In: Rubin MG, editor, Elsevier Inc.; 2006. p. 137-69.
34. Baumann L. Chemical Peeling. *Cosmetic dermatology. Principles and Practice*. In: Baumann L, editor. 1<sup>st</sup> ed, New York: The McGraw- Hill Companies; 2002. p. 173-86.
35. Moy LS. Superficial chemical peels with alpha-hydroxy acids. *Atlas of cutaneous surgery*. In: Robinson JK, Arndt KA, Le Boit PE, Wintroub BU, editors. Philadelphia: WB Saunders

Khunger N: Standard guidelines of care for chemical peels

- Company; 1996. p. 345-350.
36. Coleman WP 3rd, Futrell JM. The glycolic, trichloroacetic acid peel. *J Dermatol Surg Oncol* 1994;20:76-80.
  37. Monheit GD. The Jessner's + TCA peel: A medium-depth chemical peel. *J Dermatol Surg Oncol* 1989;15:953-63.
  38. Monheit GD. Combinations of therapy. Chemical peels. In: Rubin MG, editor, *Chemical Peels. Procedures in cosmetic dermatology*. Elsevier Inc.; 2006. p. 115-36.
  39. Fulton JE, Rahimi D, Helton P, Dahlberg K. Neck rejuvenation by combining Jessner/TCA peel, dermasanding and CO2 laser resurfacing. *Dermatol Surg* 1999;25:745-50.
  40. Landau M. Combination of chemical peelings with botulinum toxin injections and dermal fillers. *J Cosmet Dermatol* 2006;5:121-6.

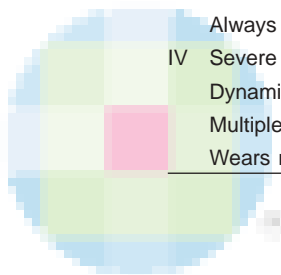
## Appendix

**Table 1: Fitzpatrick skin phototypes**

Skin type	Reaction to skin	Description
I	Always burns, never tans	Very white and freckled
II	Always burns, minimally tans	White
III	Moderately burns, uniformly tans	Light brown
IV	Minimally burns, always tans	Moderate brown
V	Rarely burns, profusely tans	Dark brown
VI	Never burns, deeply pigmented	Black skin

**Table 2: Glogau photoaging classification**

I	Mild (age 28-35 years) Little wrinkling, no scarring No keratoses Requires little or no makeup
II	Moderate (age 35-50 years) Early wrinkling, minimal scarring Shallow color with early actinic keratoses Little makeup
III	Advanced (age 50-60 years) Persistent wrinkling at rest, moderate acne scarring Discoloration with telangiectasia and actinic keratoses Always wears makeup
IV	Severe (age 60-75 years) Dynamic and gravitational wrinkling, severe acne scarring Multiple actinic keratoses Wears makeup with poor coverage



### Author Help: Online Submission of the Manuscripts

Articles can be submitted online from <http://www.journalonweb.com>. For online submission articles should be prepared in two files (first page file and article file). Images should be submitted separately.

- 1) **First Page File:**  
Prepare the title page, covering letter, acknowledgement, etc., using a word processor program. All information which can reveal your identity should be here. Use text/rtf/doc/pdf files. Do not zip the files.
- 2) **Article file:**  
The main text of the article, beginning from Abstract till References (including tables) should be in this file. Do not include any information (such as acknowledgement, your names in page headers, etc.) in this file. Use text/rtf/doc/pdf files. Do not zip the files. Limit the file size to 400 kb. Do not incorporate images in the file. If file size is large, graphs can be submitted as images separately without incorporating them in the article file to reduce the size of the file.
- 3) **Images:**  
Submit good quality color images. Each image should be less than **1024 kb (1 MB)** in size. Size of the image can be reduced by decreasing the actual height and width of the images (keep up to about 6 inches and up to about 1200 pixels) or by reducing the quality of image. JPEG is the most suitable file format. The image quality should be good enough to judge the scientific value of the image. Always retain a good quality, high resolution image for print purpose. This high resolution image should be sent to the editorial office at the time of sending a revised article.
- 4) **Legends:**  
Legends for the figures/images should be included at the end of the article file.