Treatment of facial burn sequelae using fractional CO\textsubscript{2} lasers in patients with skin phototypes III to VI

Tratamento de sequelas de queimadura de face com laser de CO\textsubscript{2} fracionado em pacientes com fototipos III a VI

ABSTRACT

Background: Reports on improvement in post-traumatic or pathological scars with the use of fractional CO\textsubscript{2} laser (CO\textsubscript{2}F) conclude that it is a safe and effective technology, though used only in patients with phototypes II to III. The aim of this study was to evaluate the effectiveness of CO\textsubscript{2}F in patients with facial burn sequelae with phototypes III to VI. Methods: A total of 14 patients (average age, 29 years) with facial burn sequelae and phototypes III to VI were subjected to a CO\textsubscript{2}F laser treatment. After 2 months, the burns were evaluated using a 6-parameter scale, including color, texture, hydration, surface irregularities, volume, and distensibility. Results: The average durations of pain, edema, and hyperemia were 19 hours, 1.3 days, and 6.5 days, respectively. The fall of crusts was completed between 5 and 36 days with an average of 13.4 days. Two months after the session, 5 patients developed punctiform hypochromia in a checkerboard pattern corresponding to the points of laser ablation. The subjective satisfaction of the evaluators (i.e., both patients and physicians) with the treatment was 84.6%. The patients reported improvements in surface irregularities, distensibility, and skin texture (57% of the cases); hydration (43%); volume (28%); and color (14%). Meanwhile, the doctors reported improvements in surface irregularities and distensibility (43%). Conclusions: The treatment with CO\textsubscript{2}F laser with mild parameters was well tolerated and resulted in high satisfaction rates for patients with facial burn sequelae as well as improved skin texture, distensibility, and surface irregularities. The high incidence of hypopigmentation must be considered while prescribing CO\textsubscript{2}F laser treatment to patients with phototypes IV to VI.

Keywords: Burns/complications. Carbon dioxide. Laser therapy/methods.
Resultados: A duração média da dor foi de 19 horas; do edema, 1,3 dia; e da hiperemia, 6,5 dias. A queda das crostas finalizou entre 5 dias e 36 dias, com média de 13,4 dias. Dois meses após a sessão, 5 pacientes evoluíram com hipocromia puntiforme no padrão quadriculado correspondente aos pontos de ablação do laser. A satisfação subjetiva dos avaliadores (pacientes e médicos) com o tratamento foi de 84,6%. Para os pacientes, houve melhora das irregularidades de superfície, da distensibilidade e da textura da pele (57% dos casos), da hidratação (43%), do volume (28%) e da cor (14%). Para os médicos, houve melhora das irregularidades de superfície e da distensibilidade (43%). Conclusões: O tratamento com laser de CO₂F com parâmetros suaves foi bem tolerado e apresentou alto índice de satisfação em pacientes com queimeadura facial, com melhora da textura, distensibilidade e irregularidades de superfície. A alta incidência de hipopigmentação é um fator a ser considerado na indicação do laser de CO₂F em pacientes com fototipos IV a VI.


INTRODUCTION

Whether restored spontaneously or grafted, the skin exhibits permanent changes after burns, especially a reduction in the number of appendages. Changes in collagen and elastic fibers as well as differences in the composition of the extracellular matrix lead to a loss of skin mechanical properties; clinically, this means reduced distensibility, especially in hypertrophic scars1.

The improvement of the quality of post-burn skin, which is usually dry, scaly, dyschromic, and inelastic, can help improve patients’ self-esteem and quality of life2.

In recent last years, fractional ablative lasers, especially fractional (CO₂F) carbon dioxide (CO₂) lasers, have gained acceptance as the preferred method for skin resurfacing. They minimize wrinkles, photoaging, acne scars, and sagging skin without the disadvantages or risks associated with prolonged epithelialization after use of traditional ablative lasers, such as prolonged erythema, hypopigmentation, and even scars. With CO₂F lasers, a large part of the skin remains intact around the microablation columns3.

Recently, reports on the improvement of non-acne scars, including keloids, using CO₂F lasers have appeared in the literature4-6. In atrophic scars, CO₂F lasers reduced the depth of scars by 35.6%—that is, they improved the “volume” of surface irregularities1. In 2 cases of burn sequelae, the use of CO₂F lasers resulted in contracture relaxation and improved skin surface irregularities, texture, and color4. In post-traumatic and pathological scars, CO₂F lasers are reported to improve texture, tone, and skin appearance with a low incidence of dyschromia; the authors conclude that it is a safe and effective technology for the treatment of scars, although the method has been used only in patients with skin phototypes II and III4.

Despite the existence of studies on the use of CO₂F lasers in Asian people4, there is no report on their use in patients with phototypes V and VI. However, the product manuals for laser devices present parameters for use in these patients. Furthermore, there is no existing report on the use of the Smartxide Dot® (Deka, Florence, Italy) device, which is commonly used by physicians for treating scars or burn sequelae.

Therefore, the aim of this study was to evaluate the effectiveness of CO₂F lasers in treating patients with facial burn sequelae with phototypes III to VI and determine parameters for safe use in these patients.

METHODS

Fourteen facial burn sequelae patients were selected; 12 (85.7%) were female, and 2 (14.3%) were male. Their ages ranged from 17 to 50 years (average, 29 years).

The average interval between the burn and the procedure was 19 years, ranging from 3 to 43 years.

Patients with restored tegument after second-degree burns or with grafts that exhibited dyschromia or surface irregularities were included.

All patients were followed-up at the institution for at least 3 months; they were required to use a night cream with tretinoin 0.015% to 0.05% and hydroquinone 2% to 4%, and a sunscreen with a sun protection factor of 30 or higher during the day2.

All patients signed a consent form prior to study participation.

One out of the 14 individuals included in the study did not carry out the treatment on the entire face because after performing a test in a periauricular area of 2 cm², the patient developed significant hypochromia. The remaining 13 patients were subjected to the laser on the entire area affected by burn sequelae.

The classification of the skin phototype according to Fitzpatrick ranged from III to VI (Figure 1).
**Laser Parameters**

A CO₂ F Smartxide Dot laser was used. One hour before the application, the patients received 1 g dipyrone orally.

The CO₂ F laser was applied using a cold-air blast to reduce pain sensitivity (Freddo® – Fabinject, Taubaté, SP, Brazil).

The parameters used for the session were stack 1 (single shot), 30 W power, spacing (Sp) 1.000 µm, and dwell time (DT) or exposure time of 500 µs.

**Clinical Evolution**

The following parameters were evaluated:

- a) Pain score, using a scale from 0 to 10
- b) Duration of pain
- c) Duration of edema
- d) Duration of hyperemia
- e) Time for crusts to fall off
- f) Presence of dyschromia
- g) Presence of itching or other symptoms

The initial follow-up period in this pilot study was 2 months, followed by weekly follow-ups until complete epithelialization and monthly follow-ups thereafter. The present patients will be followed-up for 1 year.

**Scar Evaluation Method**

Burns were evaluated by the patients and 3 medical staff members by using a 6-parameter scale, including color, texture, hydration, surface irregularities, volume, and distensibility.

Each parameter was scored from 0 to 3 as follows: 0 = “unsatisfactory,” 1 = “regular”, and 2 = “good.” The medical evaluators were provided with an explanatory table to guide the scoring of each parameter (Chart 1). For some parameters, extreme cases could receive a score of -1, which corresponded to “very bad.” The sum of the scores of each parameter was taken as the final score, which ranged from 0 to 12.

The medical evaluation score was obtained by averaging the scores of the 3 medical evaluators.

**RESULTS**

On a scale from 0 to 10, the average pain reported by patients during laser application was 5.2. The average duration of pain or burning was 19 hours, ranging from 3 hours in 8 patients to 2–3 days in 5 patients.

Facial edema developed in 9 patients, and persisted on an average for 1.3 days. The average time of hyperemia was 6.5 days. The crusts started to fall at 2 days, and all crusts fell between 5 and 36 days (average, 13.4 days).

Three patients reported pruritus during the phase when the crusts were falling off. One patient had an allergic reaction during this period, probably related to the sunscreen used, with good resolution after 15 days. This was the only patient who would not submit to a new session of treatment because she considered the post-treatment very uncomfortable.

Two months after the completion of the session, 5 patients developed hypochromia characterized by the presence of multiple discrete points of hypochromia in a checkerboard pattern corresponding exactly to the points of injury made by the laser. Among the patients who developed hypochromia, 2, 2, and 1 were phototype IV, V, and VI, respectively (Figure 2). One of the patients with phototype IV did not complete the session because hypochromia was observed after a test in a 2-cm² periauricular area, which is considered an area of high intensity as mentioned earlier.

The mean evaluation scores of the patients were 7 and 7.6 before and after the treatment, respectively. The average medical assessment score increased from 5.3 to 6.8. Skin surface irregularities, distensibility, and texture improved in 57% of the cases, hydration in 43%, volume in 28%, and color in 14%. The scores of the medical evaluators revealed improved skin surface irregularities and distensibility in 43% of the cases and improvement of skin texture, hydration, color, and volume in 28% of the cases. The rate of subjective satisfaction of the evaluators (i.e., both patients and physicians) with the treatment was 84.6% (Figure 3).

**DISCUSSION**

This pilot study revealed important aspects to guide future protocols of burn sequelae treatment. The subjective satisfaction rate of the patients and doctors of 84.6%, especially regarding the improvement in texture, hydration, and distensibility of the post-burn tegument and smoothing of surface irregularities, is emphasized.

However, the scale used in the present study was not sensitive enough to measure satisfaction because “some improvement” in one of the requirements may not be enough to change a score from 1 to 2.
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Chart 1 – Table used by the medical evaluators to score burn sequelae.

<table>
<thead>
<tr>
<th>Color (compared to normal skin)</th>
<th>-1 (very bad)</th>
<th>0 (unsatisfactory)</th>
<th>1 (regular)</th>
<th>2 (good)</th>
</tr>
</thead>
<tbody>
<tr>
<td>High contrast with hypo- and hyperchromic areas</td>
<td>Highly visible contrast</td>
<td>Light contrast</td>
<td>No contrast</td>
<td></td>
</tr>
</tbody>
</table>

| Texture (movement parallel to the skin) | Diseases with extreme harshness | Bad texture | Acceptable texture | Soft skin, good/excellent texture |

| Hydration | Diseases with extreme dryness | Bad | More or less | Good |

| Surface irregularities | | | | |
|------------------------|---------------------------|-------------------------------|--------------------------|
| Keloid or retractable scar with irregularities in body contouring | Very depressed scar or hypertrophic | Regular | Scar that does not cause change in body contouring |

| Distensibility (elasticity, perpendicular traction to the skin surface) | No distensibility, skin adhered to the bone plane | Bad, distensibility, markedly impaired | Regular, distensibility moderately worsened related to normal skin | Response similar to the normal skin traction around |

Figure 2 – Patient with skin phototype VI, with points of hyperchromia after treatment.

Figure 3 – A 23-year-old patient before and 2 months after fractional CO₂ laser treatment.

The parameters listed in the manual for the use of the laser on scars and keloids are 30 W power, Sp 600 µm (30% of injury with 70% of skin preserved), and DT ranging from 1.000 µs (phototype I) to 500 µs (phototype V). These parameters were considered to be very aggressive during the design of this study; thus, we opted to widen the Sp to 1.000 µm (ablation of 10% of the treated area) and use a DT of 500 µs (less deep injury); this turned out to be effective. The epithelialization time of 13.4 days is higher than those observed for rejuvenating treatments or acne sequelae; this is due to the smaller number of skin appendages in burn sequelae.

Even though the parameters of ablation were considered to be mild, 35.7% of cases exhibited hypochromia. A previous study reports a hyperchromia rate of 11.1% 6 months after CO₂F laser application.

The parameters of ablation were considered to be mild, 35.7% of cases exhibited hypochromia. A previous study reports a hyperchromia rate of 11.1% 6 months after CO₂F laser application.

The patients in the present study will be followed-up for 1 year with topical treatment. The late evolution will determine whether the treatment with the CO₂F laser is in fact safe as postulated.

We suggest the use of even milder parameters in patients with phototypes IV to VI as well as the performance of a treatment test on a small area. Even the 4 patients who developed hypochromia expressed satisfaction with the improvement in other skin aspects, including medical evaluation. Punctiform hypochromia was considered to be very discrete by patients and more easily camouflaged than the surface irregularities.

All patients expressed a desire to be part of the full duration of this study, which includes 3 sessions.

The mechanical properties of the burn sequelae are thought to have improved because the CO₂ laser modulated...
tissue repair, resulting in increased basic fibroblast growth factor (bFGF) and suppressed secretion of transforming growth factor β1 (TGFβ1). Thus, CO₂ lasers simultaneously promote cell replication and have the potential to balance collagen organization against excessive fibrosis, preventing aberrant tissue repair¹⁰⁻¹¹. Cytokine secretion after CO₂F laser treatment was different from the secretory pattern observed after tissue injury. The presence of subsequent bFGF secretion in contrast to TGFβ1 reduction until the 30th day confirms that the injury induced by this laser is much more physiological as compared to that observed in hypertrophic scars³.

**CONCLUSIONS**

The treatment with the CO₂F laser with mild parameters was well tolerated and resulted in high satisfaction rates in patients with facial burn sequelae with apparent improvement in skin texture, distensibility, and surface irregularities. The high incidence of hypopigmentation must be considered when considering this treatment for patients with phototypes IV to VI.

**REFERENCES**