Background: Melasma is an acquired, chronic hypermelanosis condition. Melasma is more common in women of all races and occurs especially on the face. The pathogenesis of melasma is very complex and the treatment is still a challenge. The purpose of this study was to report mixed-type melasma treated with low fluence Q-switched Nd-YAG 1064 nm laser.

Case Presentation: A case of melasma in a 54-year-old woman was reported. Dermatological examination showed presence of brownish macules and patches in the centrofacial area with symmetrical distribution. Examination with a wood lamp showed mixed type. Patients have received topical therapy but there were no improvement.

Results: The patient was then treated with low fluence Q-switched Nd-YAG 1064 nm laser for three sessions with an interval of 2 weeks. At 6 weeks of treatment the modified MASI (mMASI) value was reduced from 8.4 to 4.6 and the VAS value was increased from 2 to 8.

Conclusion: The depth of the pigment determined the response to therapy. In the mixed type melasma the response for therapy is only partial. The low-fluence Q-switched Nd-YAG 1064 nm laser can penetrate deeper into the dermis and damage melanin in a short time.

Keywords: mixed-type melasma, low fluence, Q-switched Nd-YAG laser

Correspondence: Aninda Fitri Nugrahani. Department of Dermatology and Venereology, DR. Saiful Anwar Hospital, Malang, East Java/Faculty of Medicine, Universitas Brawijaya, Malang, East Java. Email: aninda16fitri@gmail.com


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chemical peeling, and various laser therapy (Choi et al, 2015). Melasma often does not improve with chemical peeling therapy or topical therapies such as hydroquinone, retinoid, azelaic acid or kojic acid due to resistance to these agents and rapid recurrence in the formation of melasma lesions (Sim et al, 2015). Since 2006, the use of low-fluence 1064-nm Q-switched Neodymium: Yttrium-Aluminum-Garnet (QS Nd-YAG), a technique known as "laser toning", became popular for melasma therapy. Although said to be effective, this Nd-YAG QS laser can cause hypopigmentation in some patients(Kim et al, 2013).

In this case report a 54-year-old woman with mixed-type melasma will be treated with a 1064-nm low-fluence Q-switched Neodymium-Yttrium-Aluminum-Garnet laser.

**CASE PRESENTATION**

A 54-year-old woman came to the Dermatology and Venereology Clinic of Dr. Saiful Anwar Hospital Malang, with complaint of dark spots on his face. These spots have appeared since 1 year ago. Spots initially appear in the nose area, then spread to the right and left cheeks and forehead. The spots do not feel itchy, hot, or painful. Before the spots appear the patient denies the presence of acne or irritation in the area. One month before going to the hospital the patient visited his son who lived in Kalimantan and the patient felt his spots darken.

Patient is a housewife. Besides doing household activity, she also actively participating in the recitation around the house. The patient said that she had routinely used cream from a beauty clinic >5 years. About 4 months ago the patient changed her beauty clinic and got a morning and night cream. The creams are used for 2.5 months. The patient said there was no change in the spots. The past month, she has not used any treatment cream. Patients said she never wear sunscreen routinely every day.

The patient has 2 children. Between the first and the second child, she used implant contraceptive for 5 years. After the birth of the second child, she underwent tubectomy. The patient said that she had never taken hormonal drugs such as birth control pills or other drugs on a regular basis. The patient said that neither the patient's mother nor the patient's brother had the same complaints as the patient.

**RESULTS**

On physical examination the patient’s blood pressure was obtained 120/80 mmHg, with a pulse of 68x/ min, a respiratory rate of 20x / min, and a temperature of 36.7°C. On dermatological examination on the face at centrofacial area, patches and macules of dark brown color with firm boundaries and irregular shapes are found. On examination with Wood's lamp, it was found that there were some spots with firm boundaries and other spots with unclear boundaries.

Figure 1 depicted dermatological examination. Figure 1 showed dark brown patches and macules with firm borders and irregular shapes at the centro facial part of the face.

Figure 2 depicted wood lamp examination. Figure 2 obtained some spots with firm boundaries and other spots with indefinite boundaries.

The patient was diagnosed with mixed-type melasma. Patients were then planned to get low-fluence laser therapy Q-Switch Neodymium Yttrium Aluminum Garnet 1064 nm every 2 weeks. During the laser therapy period patients are required to wear wide-spectrum sunscreen with SPF 30 at least 2 times per day. Patients provided written informed consent to participate in the study.
Figure 1 A, B, C. Dermatological examination.

Figure 2 A, B, C. Wood lamp examination

Figure 3. Follow up of therapy. A. 4th week B. 2nd week C. 4th week D. 6th week
Before laser therapy, the patient is given topical anesthesia with 2.5% lidocaine and 2.5% prilocaine for 30-45 minutes. After that topical anesthesia was cleaned with sterile gauze moistened with 0.9% NaCl. Patients received QS: Nd-YAG laser therapy with a wavelength of 1064 nm, spot size of 8 mm, fluence 2.0 J / cm², and 3 times fit the entire face. The end point of this therapy is the presence of erythema on the skin and warm palpable skin. After therapy, a cold compress with 0.9% NaCl is applied to reduce the patient’s burning sensation. Laser therapy is performed every 2 weeks.

At the time of initial examination, the modified MASI (mMASI) score was 8.4. Two weeks after the first laser, the mMASI score decreased by 2.2, so the score on the second week became 6.2. In the fourth week the score dropped again to 4.6 and in the sixth week the mMASI score remained 4.6 (Figure 4).

Figure 4. The improvement of mMASI value

Figure 5. The improvement of VAS values
To evaluate patient satisfaction, Visual Analogue Scale (VAS) is used in the range 0-10, where 0 is no improvement and 10 is a very satisfying improvement. Evaluation was carried out at weeks 0, 2, 4, 6, and 8. VAS values obtained were 2, 5, 6, and 8. There were no significant side effects in patients (Figure 5).

**DISCUSSION**

Melasma is a chronic condition of hypermelanosis obtained on the skin, which is characterized by brown macules with irregular shapes and symmetrically distributed in areas of the body exposed to sunlight, especially the face (Handel et al, 2014). This disorder occurs most commonly between the ages of 30 and 40 years. Family history affects the occurrence of melasma in people with Fitzpatrick skin types II and III, whereas in darker skin types family history has no effect. Melasma is more common in female with a ratio of 9: 1 compared to male. In Southeast Asia the prevalence in adult women reaches 40% and in adult men 20% (Cestari et al, 2017).

Several studies have shown that there are increasing in both melanocytosis and melanogenesis in melasma patients. Some factors that can be triggers include sun exposure, use of oral contraceptives, thyroid disease, pregnancy, and medications (Sarkar et al, 2014). Study conducted by KrupaShankar et al in India shows that the most dominant trigger factor for melasma is sun exposure(KrupaShankar et al, 2014).

Facial melasma can be divided into 3 patterns based on the affected face area, namely centro facial pattern, malar pattern, and mandible pattern. In the centro facial pattern the affected area is the forehead, cheeks, upper lip, nose, and chin. Whereas in malar pattern the lesions are only found on the cheeks and nose, and in the pattern of the mandible the area involved is the ramus of the mandible(Rigopoulos et al, 2017). While based on the depth of the pigment, the type of melasma is divided into three, namely the dermal type, epidermal type, and mixed type. On Wood’s lamp examination, dermal-type melasma is clearer and darker, while in the dermal type less fluorescence can be captured. The depth of the pigmentation also affects melasma therapy. In epidermal type, the response to therapy is better than the dermal type. And in mixed types, the response produced is only partial (Sarkar et al, 2014). In this case report found melasma in 54 years old woman. Patients never use sunscreen. Dermatological examination revealed brownish macules in the centro facial area and Wood’s lamp examination showed mixed type.

The treatment of melasma consists of medical and non-medical therapy. In medical management can be given topical drugs, systemic drugs, and minimally and invasive treatment such as lasers and light and chemical peeling. Non-medical management can be topical such as skin care products, decorative cosmetics, or skin lightening cosmetics, and can also be systemic for example consumption of antioxidants (Wasitaadmaja et al, 2018).

Laser therapy in pigmented lesions is based on the theory of selective photothermolysis proposed by Anderson and Parrish. The theory states that when specific energy wavelengths are released in a shorter period of time than the thermal relaxation time (TRT) of the target chromophore, the energy will be retained in the target cell and cause less damage to the surrounding area (Arora et al, 2012). Laser quality-switched (Q-switched) produces high-intensity laser beams with very short pulse durations. The new variant of Q-switched laser is known as low fluence or Q-switched sub-thermolytic therapy(Trivedi et al, 2017). Fluence used in
sub-temolytic laser therapy is <5 J/cm² (Wattanakrai et al., 2010). Low fluence therapy using a wavelength of 1064 nm (QS: Nd-YAG) can penetrate deeper into the dermis and a little affects the epidermis (Trivedi et al., 2017).

The selection of the right beam diameter or spot size is important in the treatment of pigmented lesions. Large spot sizes are safer for the epidermis and are preferred for dark skin types (Small et al, 2016). Spot sizes between 7-10 mm produce maximum penetration to the dermis tissue. Deeper penetration will not be achieved if the spot size is above 10-12 mm (Nouri, 2014).

Patients were given a 1064 nm Q-switched Nd-YAG laser therapy with 2 J/cm² fluence, 8 mm spot size, and 3 passes. 3 laser sessions with a 2-week interval showed improvement in the lesion marked by a decrease in the mMASI score and an increase in the VAS score. There were no laser side effects in patients, either hyperpigmentation, hypopigmentation, or the presence of scars.

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Both authors developed the concept for this manuscript and edited multiple drafts of the manuscript.

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REFERENCE


