

**Evaluating the Tolerability and Effectiveness of a Biocellulose Mask following a  
Full Face Fraxel Laser Treatment  
Clinical Trial Inclusion & Exclusion Criteria  
Please Read Carefully**

**Inclusion Criteria**

- Healthy females ONLY between 35 – 60 years of age with moderate hyperpigmentation
- Subjects who do not have skin disorders
- Subjects who are willing to avoid extended periods of sun exposure for the duration of the study, including artificial tanning
- Subjects who are willing to discontinue use of any additional facial cleansers, moisturizers, and sunscreens that are not provided by the study staff
- Subjects who agree to not use any new cosmetic or toiletry products during the study
- Subjects who are generally in good health, with no evidence of acute or chronic disease or skin disorder

**Exclusion Criteria**

- Subjects who are pregnant, planning to become pregnant or breastfeeding
- Subjects with self-perceived sensitive skin
- Subjects receiving treatment within one week prior to study with:
  - Sympathomimetics (ephedrine, amphetamines)
  - Antihistamines (i.e. Claritin, Allegra, Benadryl etc.)
  - vasoconstrictors (i.e. epinephrine)
  - Non-steroidal anti-inflammatory agents (i.e. Motrin, Advil etc.)
  - Immunosuppressant's (i.e. Prednisone, Cyclosporine etc.)
  - Antibiotics and/or systemic or topical corticosteroids (Prednisone, Clobetasol etc.)
- Subjects with a history of acute or chronic skin, medical and/or physical conditions that would interfere with study material(s) and/or could influence the outcome of the study; including tattoos, piercing and sunburned skin on the face
- Immunosuppressed subjects (i.e. HIV infection, large doses of steroids)
- Subjects with a cardiac (heart) disorder
- Subjects who have used oral retinoid (Accutane® or Soriatane®) within 6 months of starting the study
- Subjects with known allergies to soaps, cosmetic and toiletry products, topical medications, sunscreens, glycolic acid or lidocaine
- Subjects with sunburn, rashes, scratches or burn marks in the treated area
- Subjects have had a facial peel, laser treatment or any cosmetic procedure within one month of starting the study
- Subjects that have used a waxing product, tanning product, tanning bed within one (1) week prior to the start of the study
- Subjects that have undergone chemotherapy or radiation
- Subjects currently participating or planning to participate in any other clinical trial during the course of the study