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**Hall et al.**

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(54) **SKIN GRAFT DEVICES AND METHODS**

2017/00561 (2013.01); A61B 2017/00734  
(2013.01); A61B 2017/306 (2013.01)

(71) Applicant: **KCI Licensing, Inc.**, San Antonio, TX  
(US)

(58) **Field of Classification Search**  
CPC ..... A61B 17/322; A61B 17/32053  
See application file for complete search history.

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U.S.C. 154(b) by 126 days.

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**Related U.S. Application Data**

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Apr. 19, 2012, now Pat. No. 9,468,459.

Louis C. Argenta, MD and Michael J. Morykwas, PHD; Vacuum-  
Assisted Closure: A New Method for Wound Control and Treat-  
ment: Clinical Experience; Annals of Plastic Surgery.

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(51) **Int. Cl.**

*Primary Examiner* — Richard Louis

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**A61B 17/3205** (2006.01)

**A61B 10/02** (2006.01)

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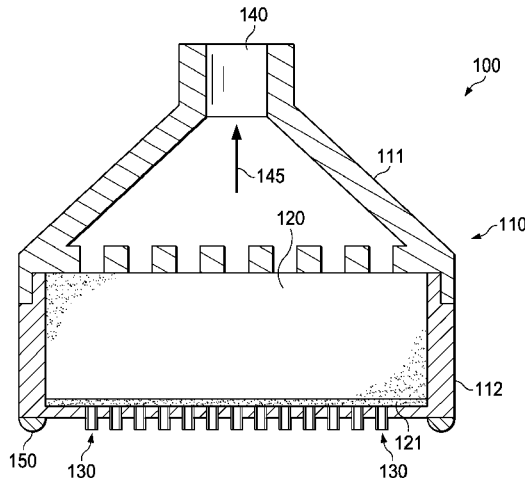
(57) **ABSTRACT**

(52) **U.S. Cl.**

Devices and methods for obtaining a plurality of skin tissue  
particles for use in skin grafting.

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(2013.01); **A61B 10/0233** (2013.01); **A61B**

**7 Claims, 18 Drawing Sheets**



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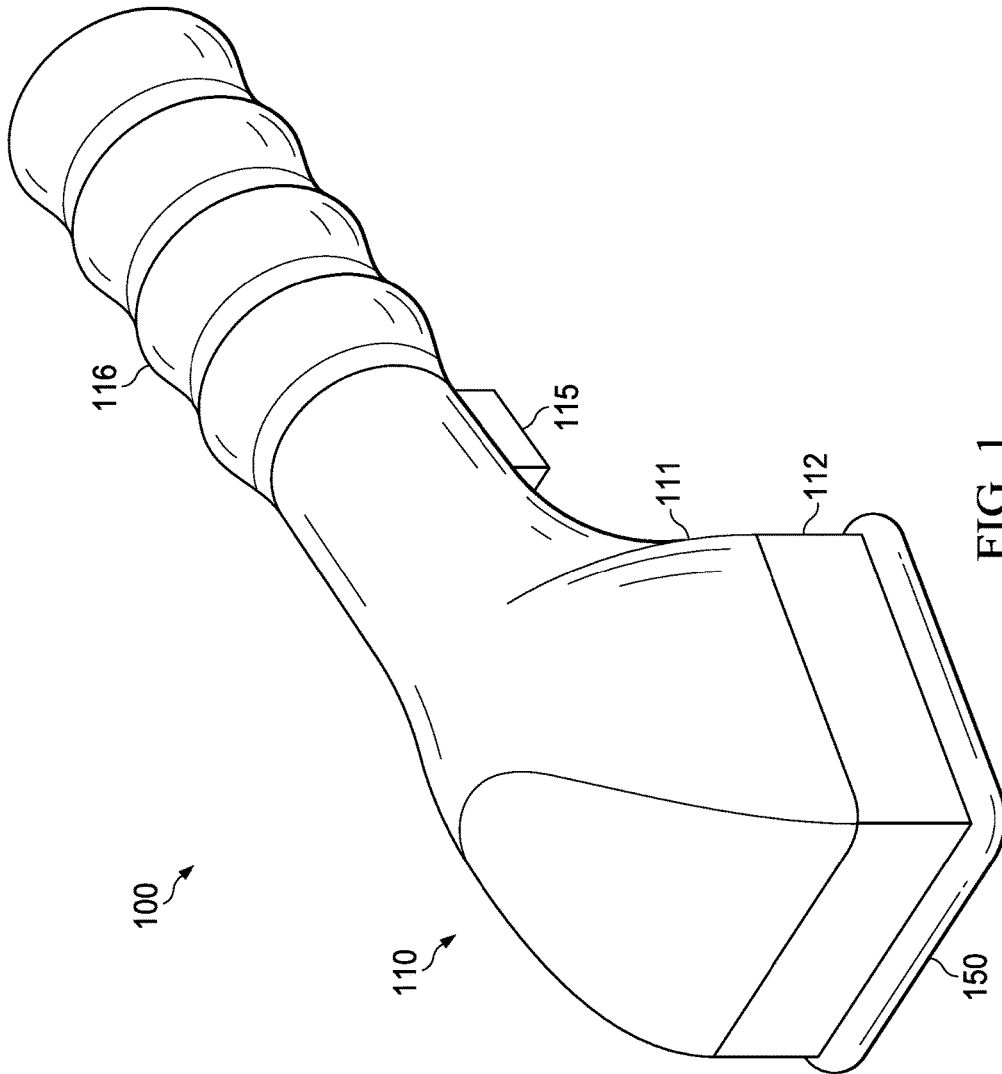
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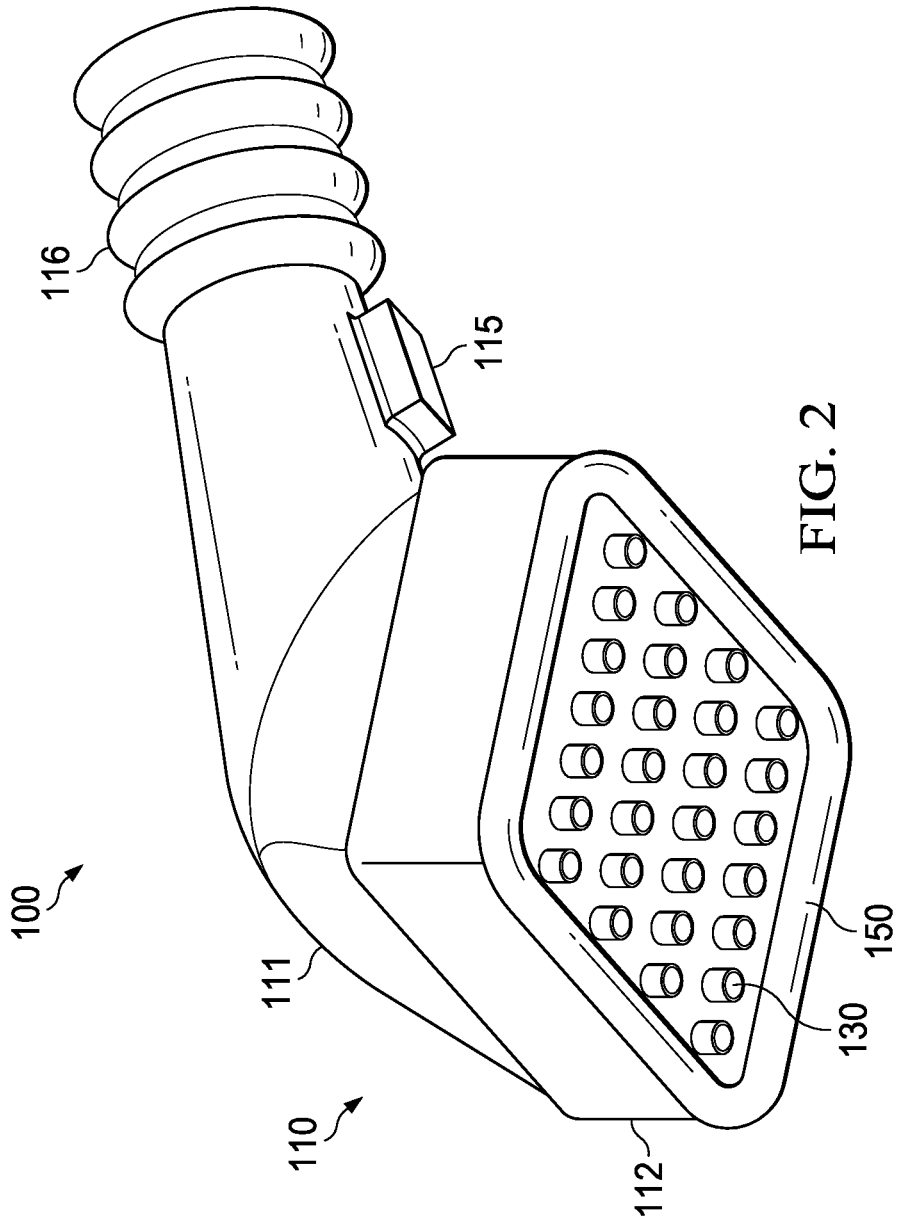
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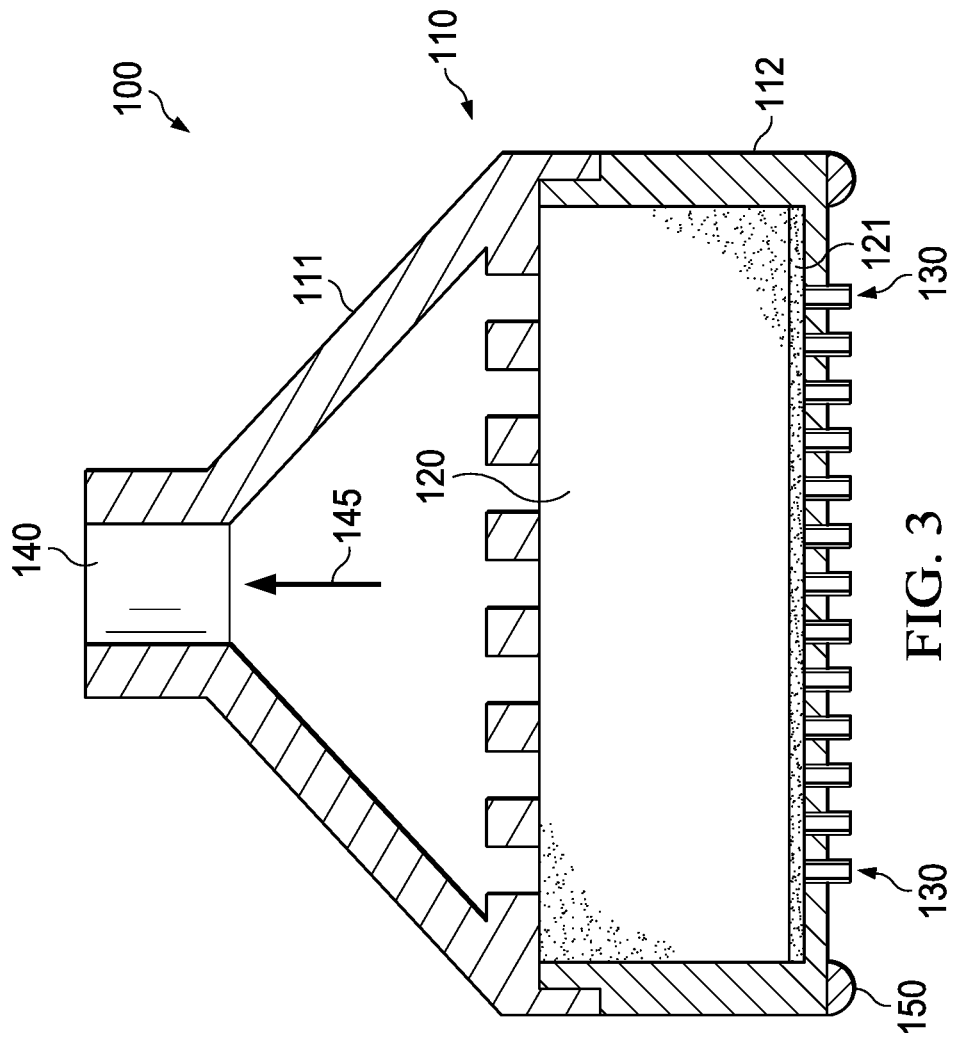
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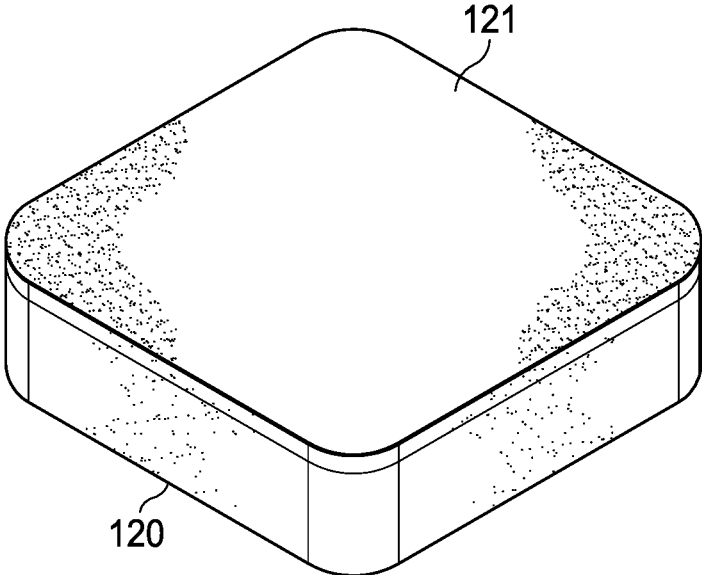


FIG. 4

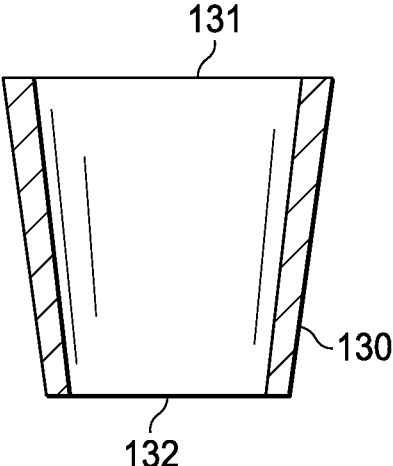


FIG. 5

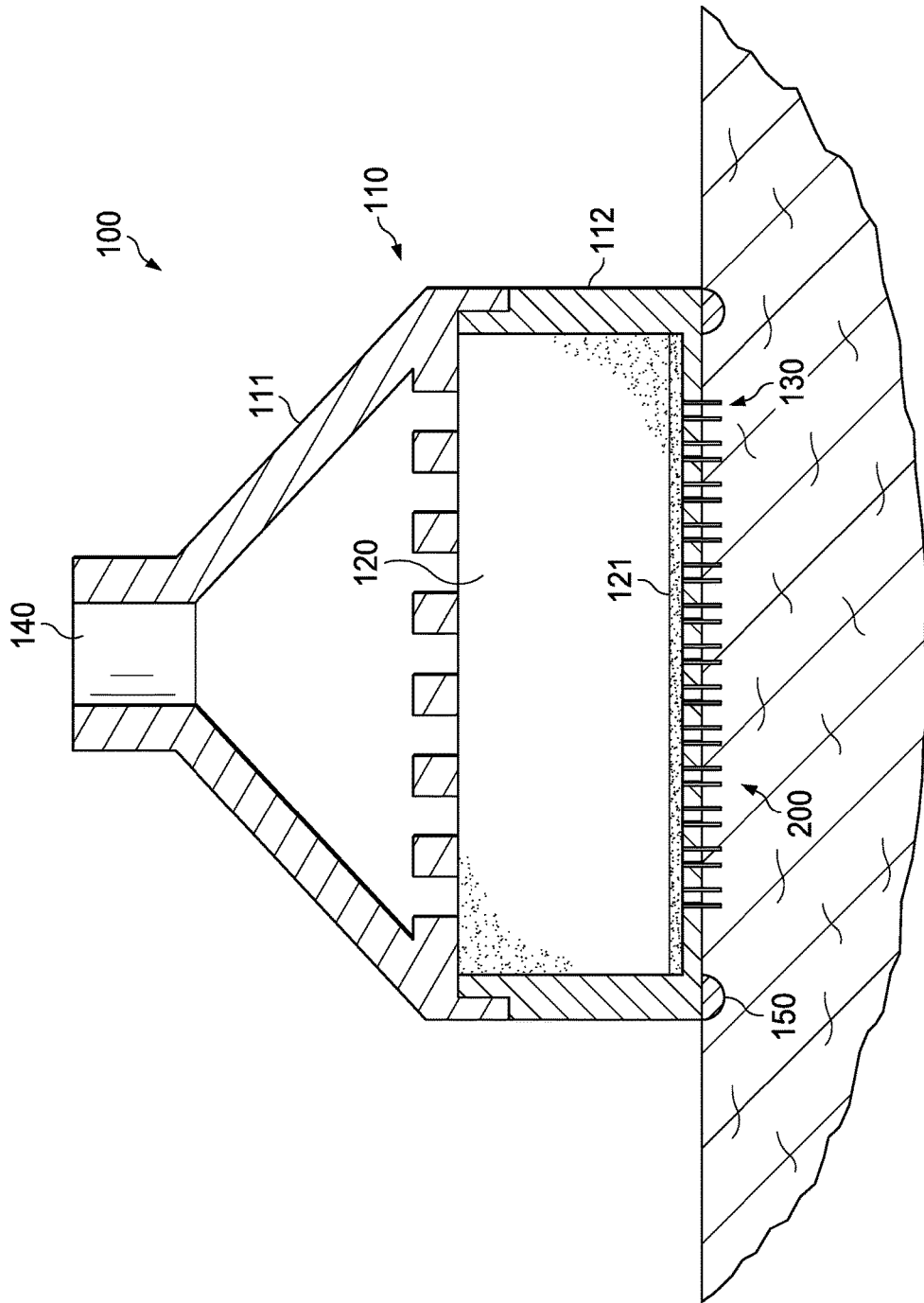


FIG. 6



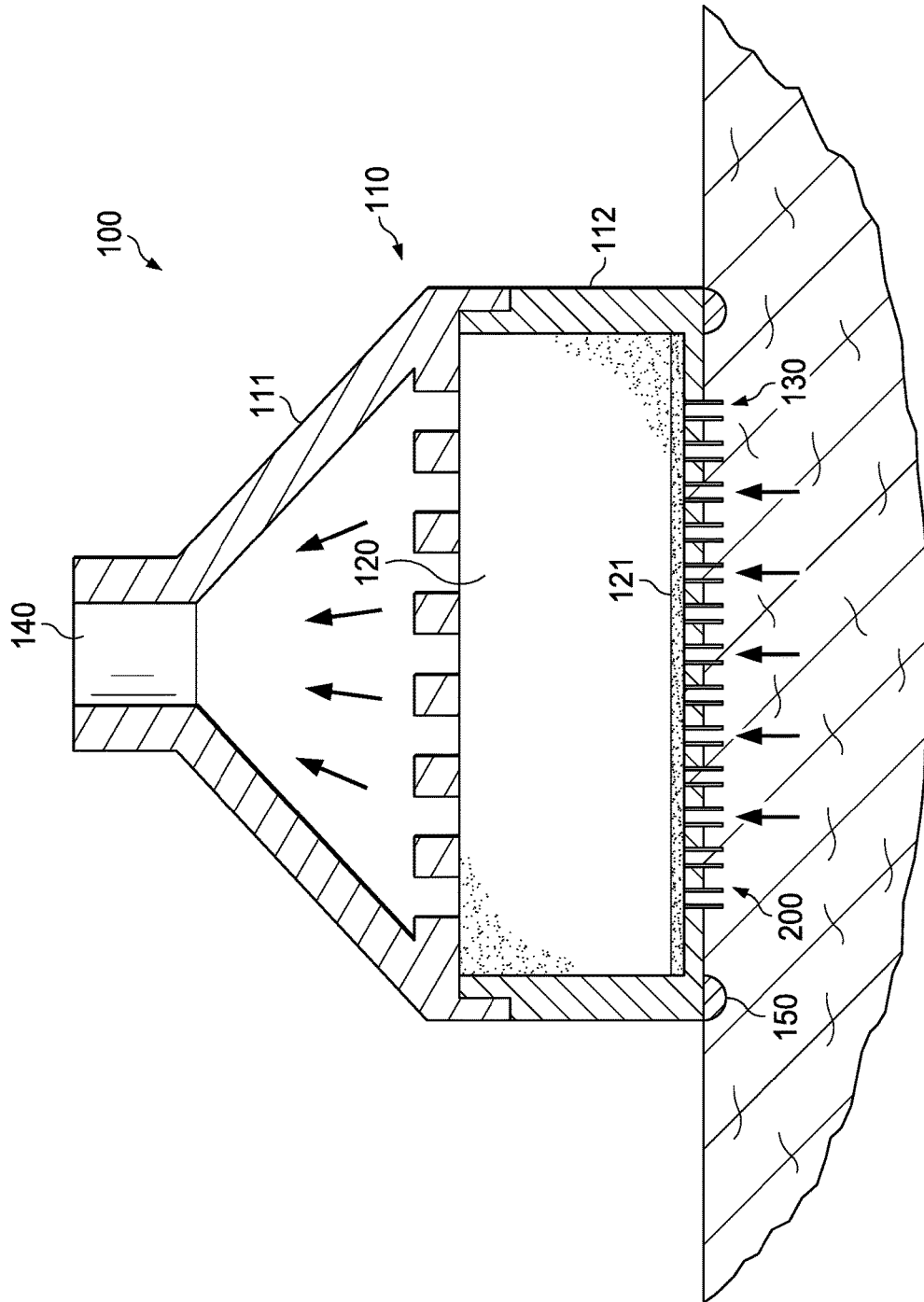


FIG. 7

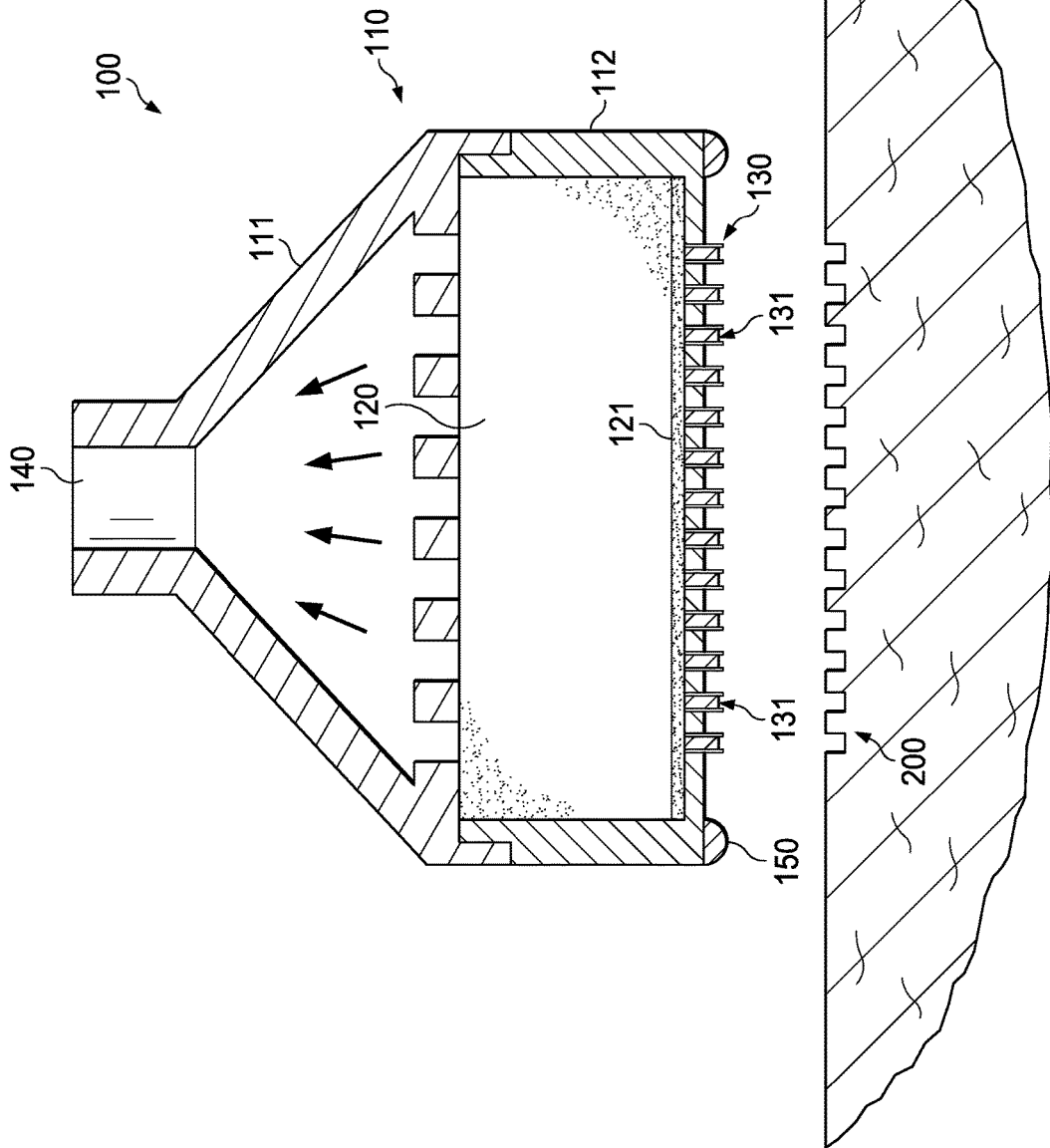
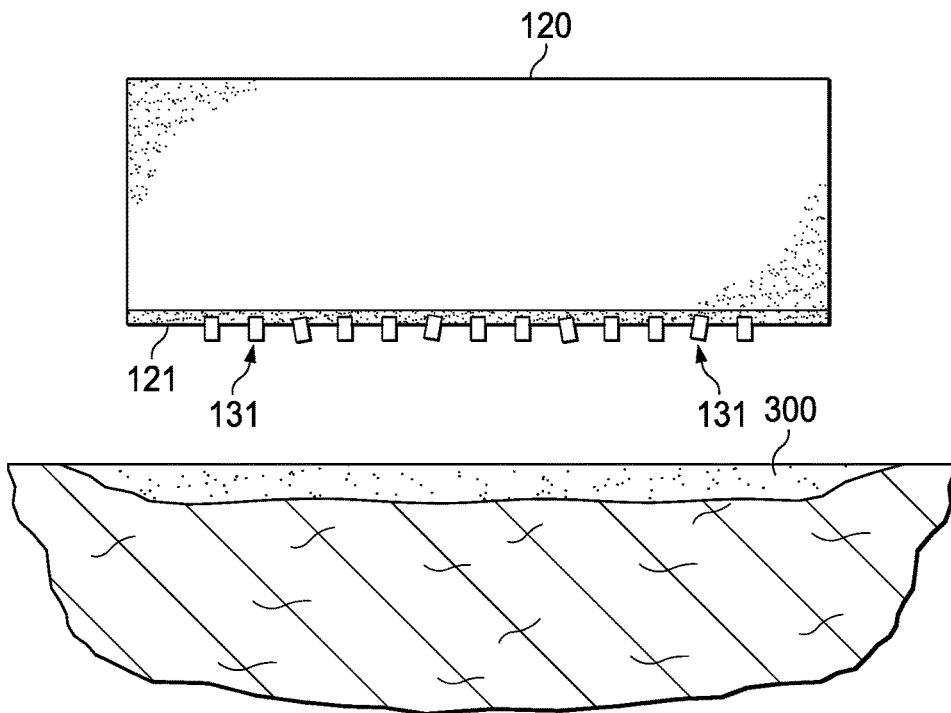
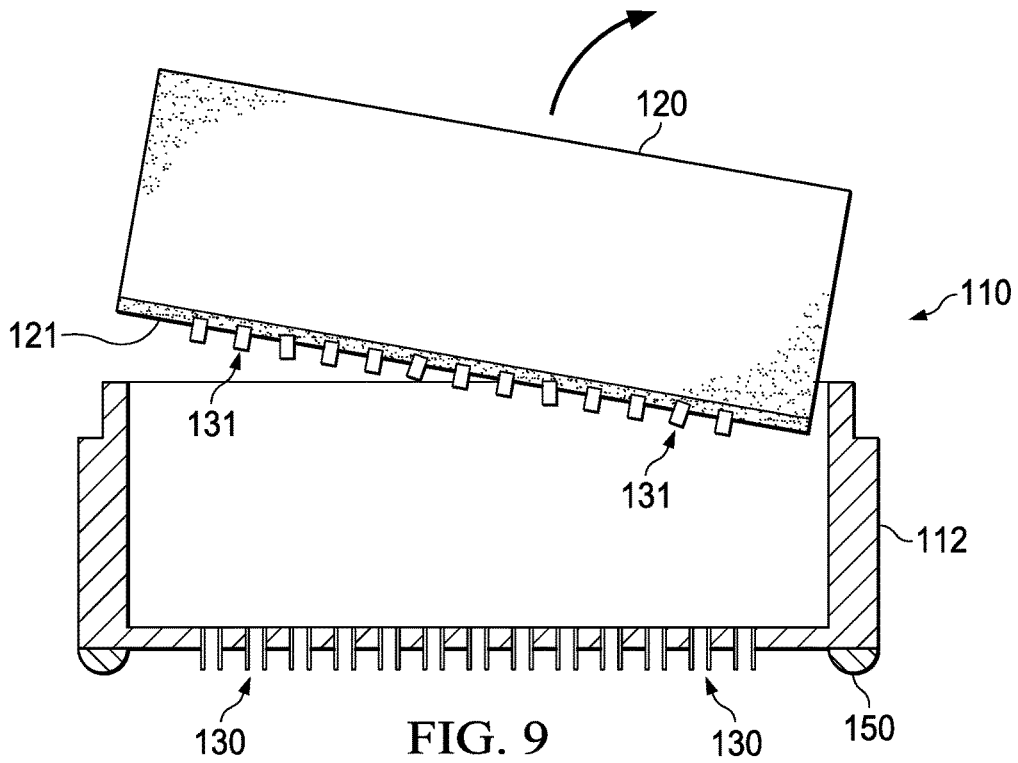


FIG. 8



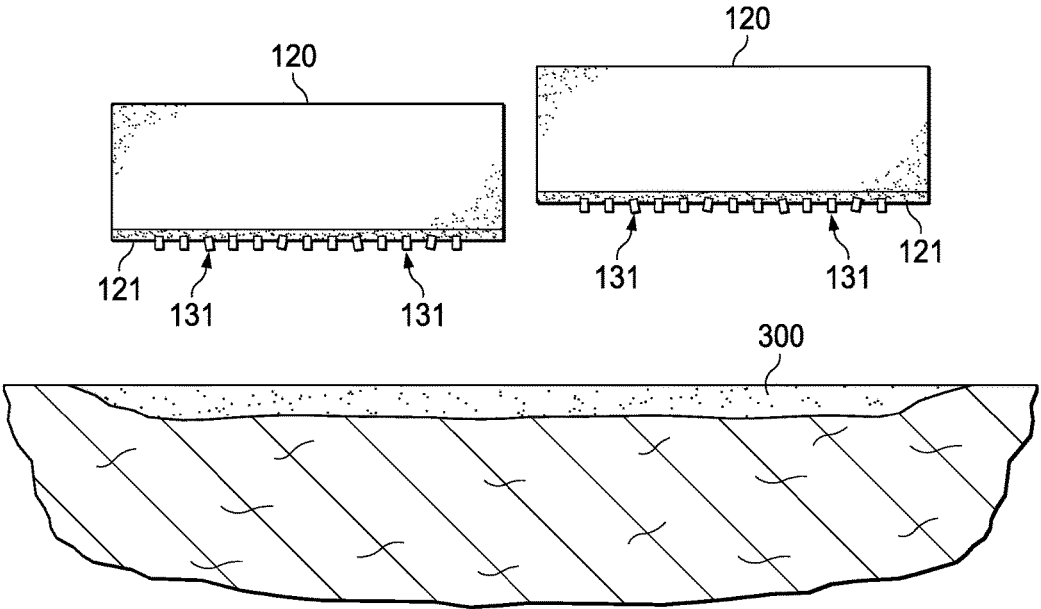


FIG. 11

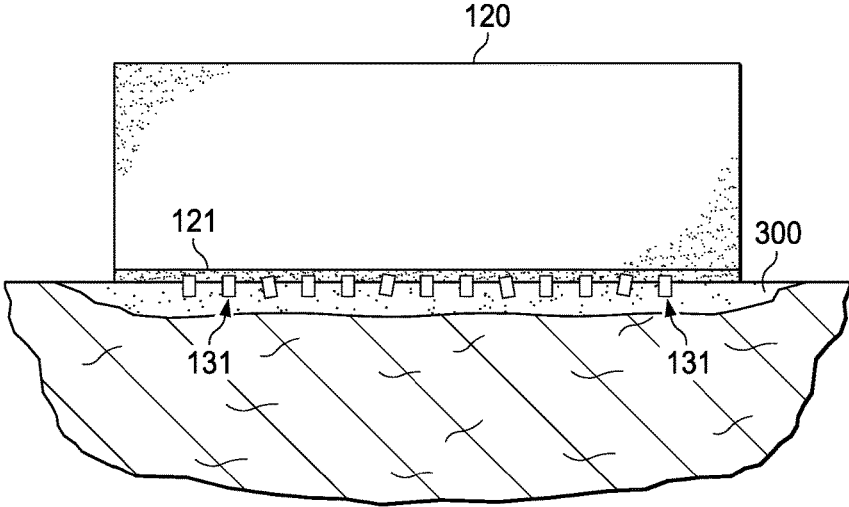


FIG. 12

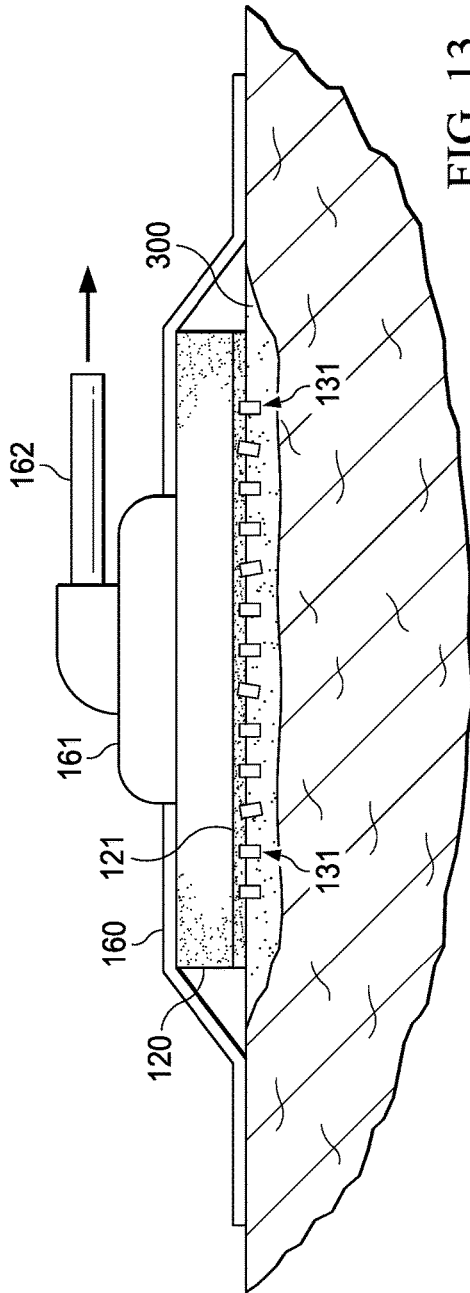


FIG. 13

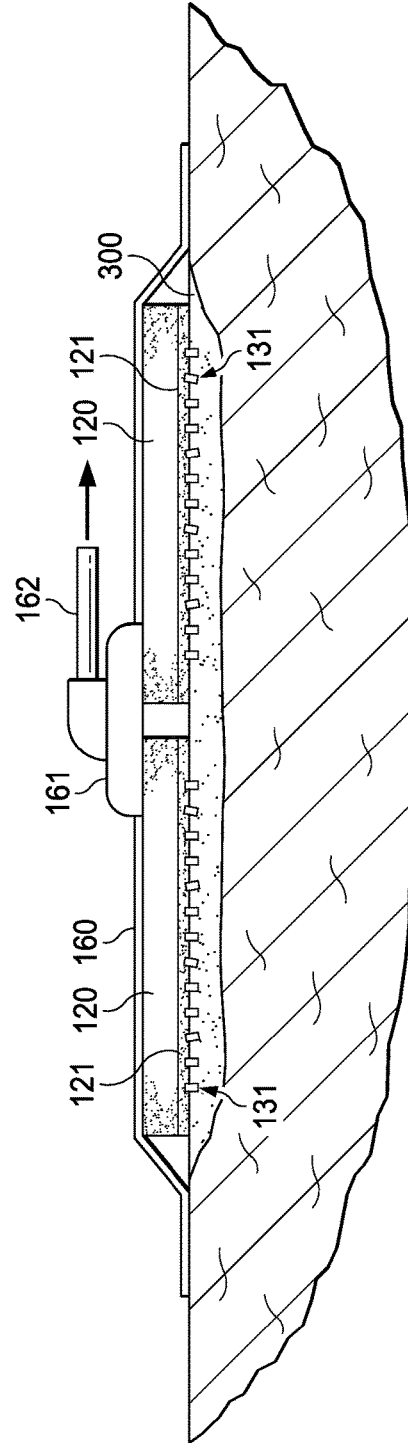
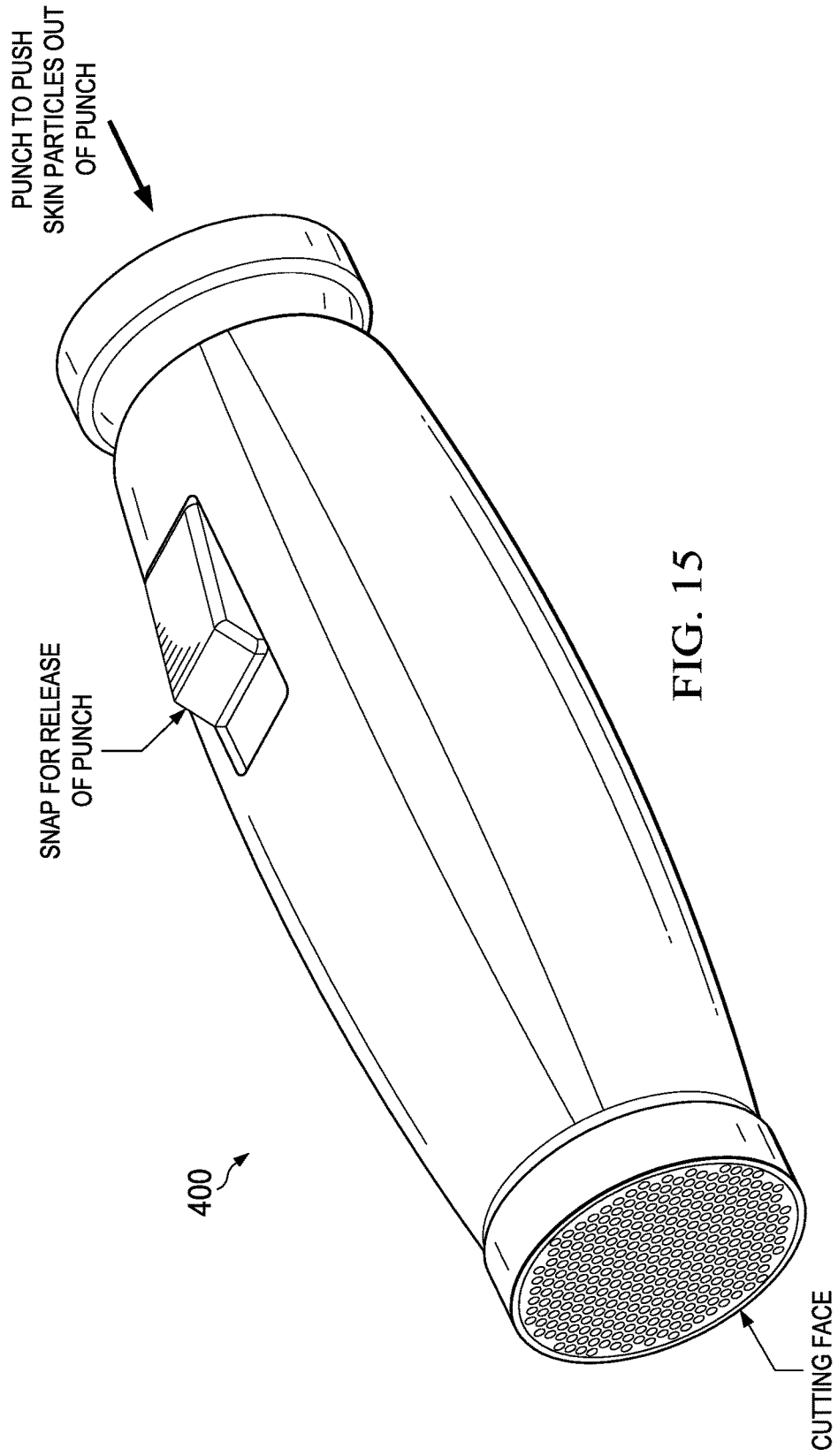


FIG. 14



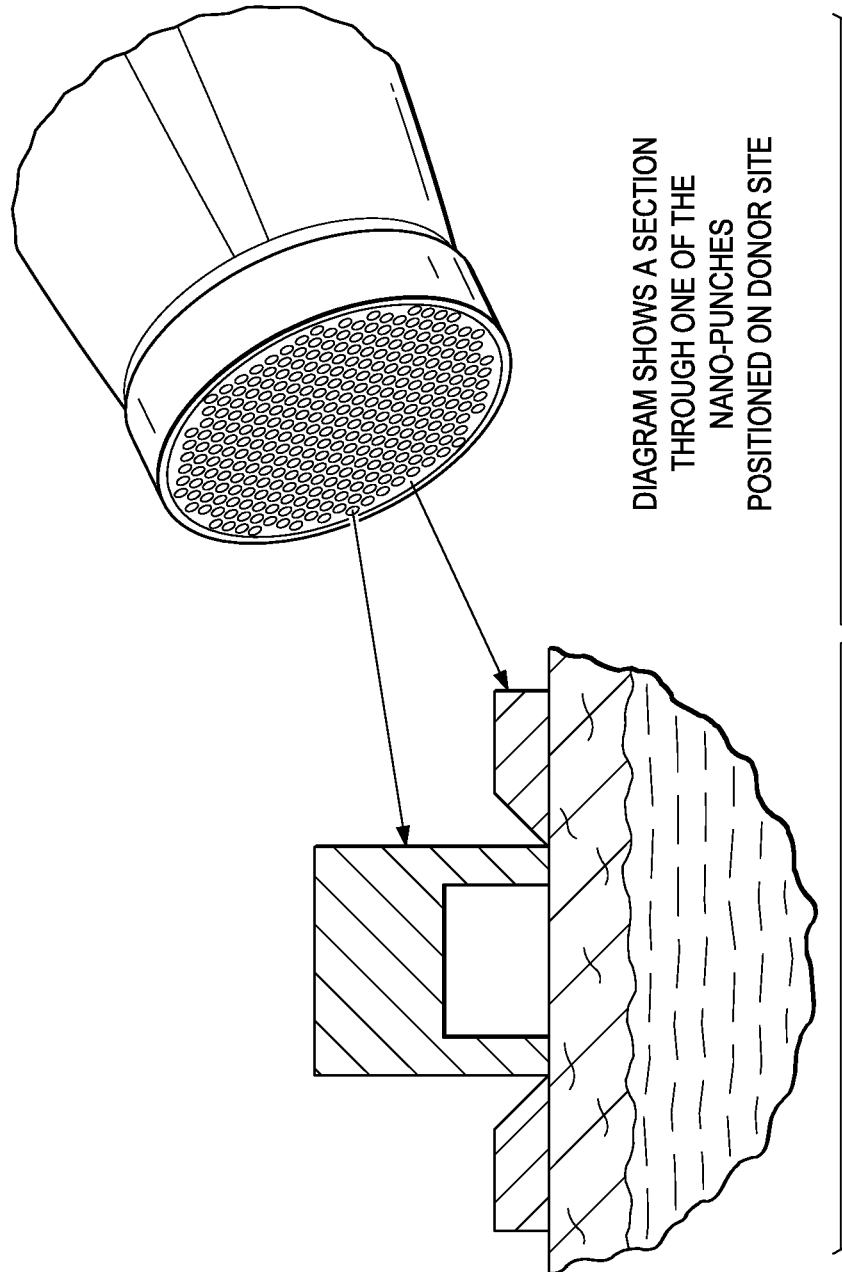


FIG. 16

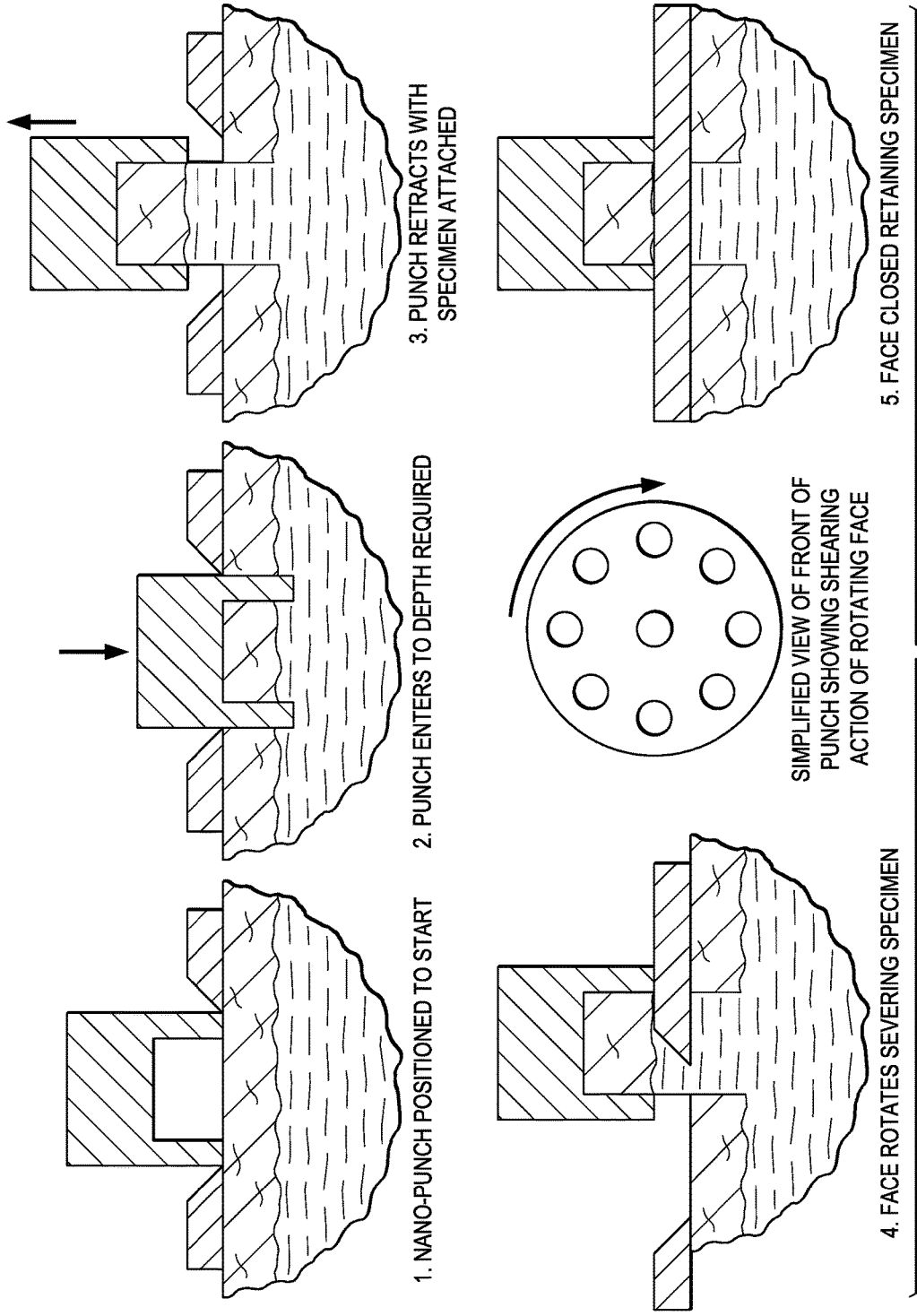


FIG. 17



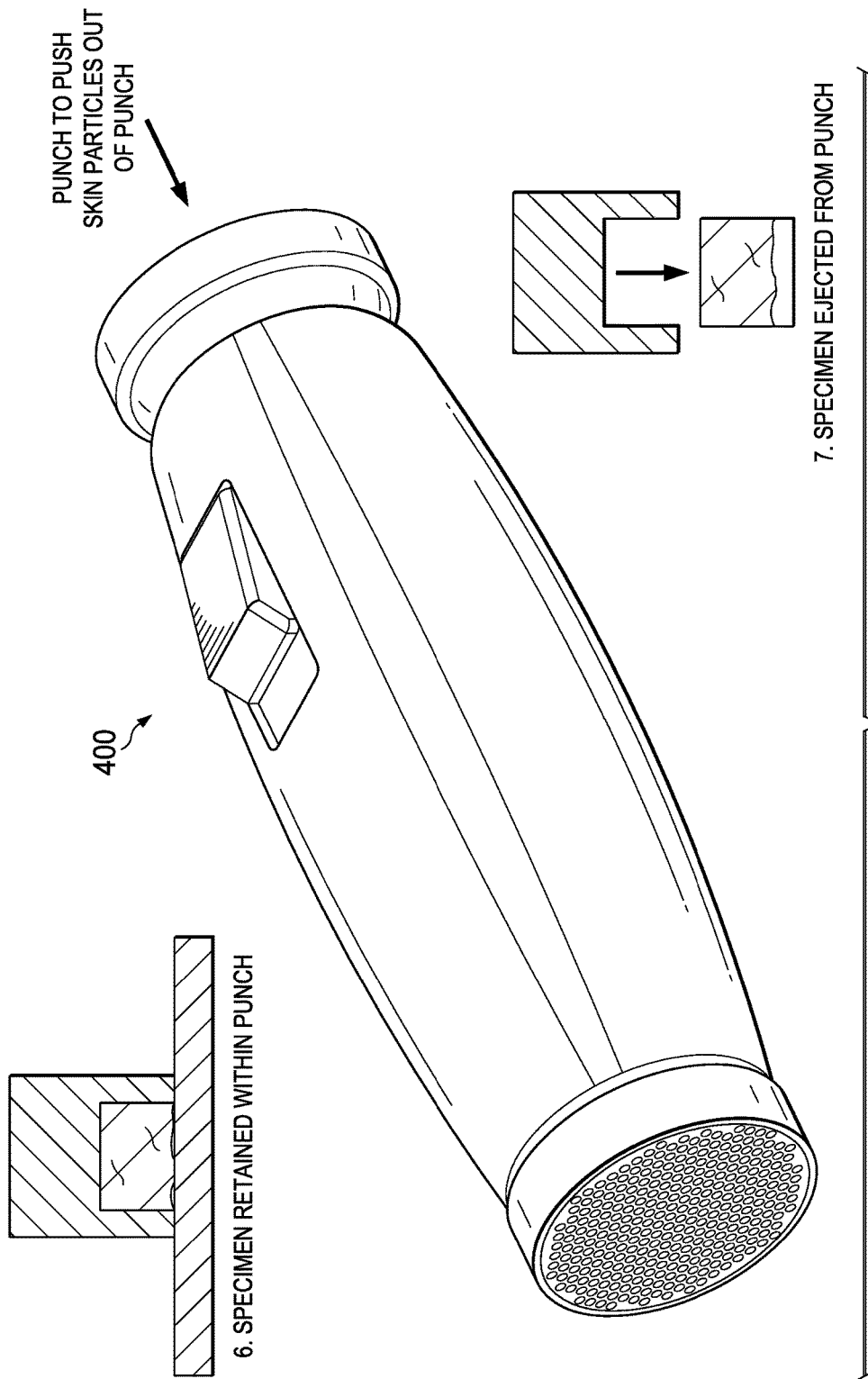


FIG. 18

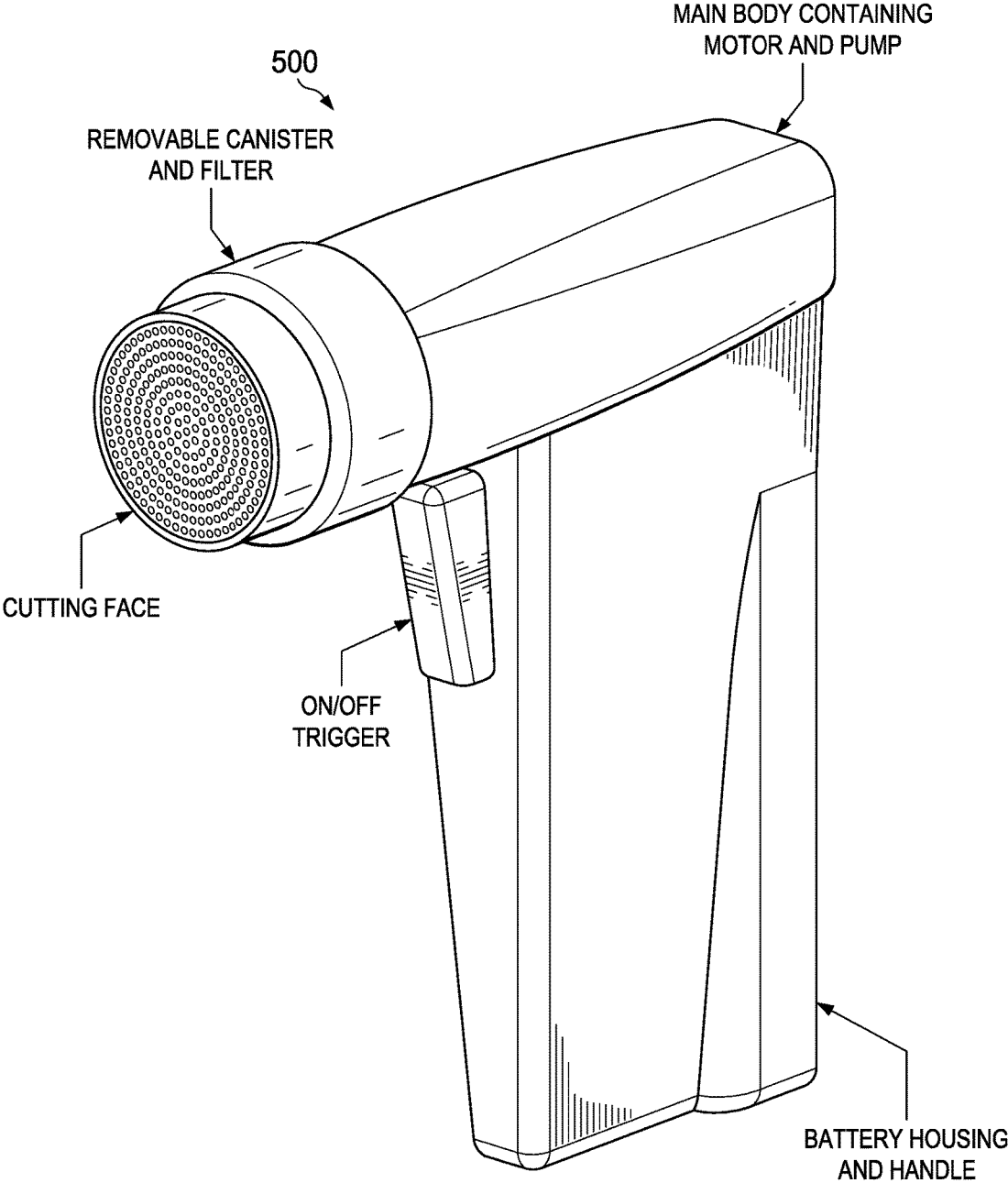


FIG. 19

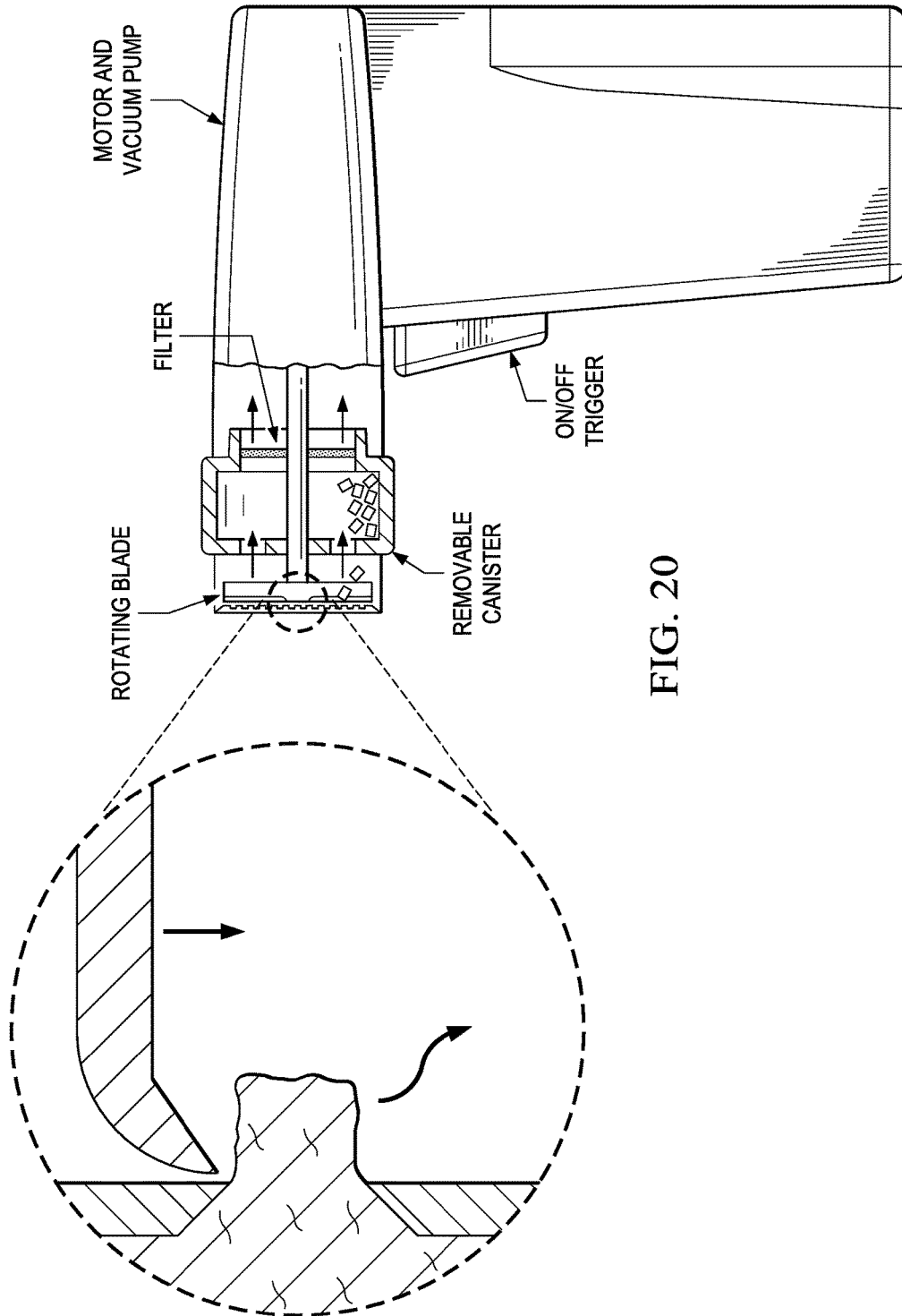
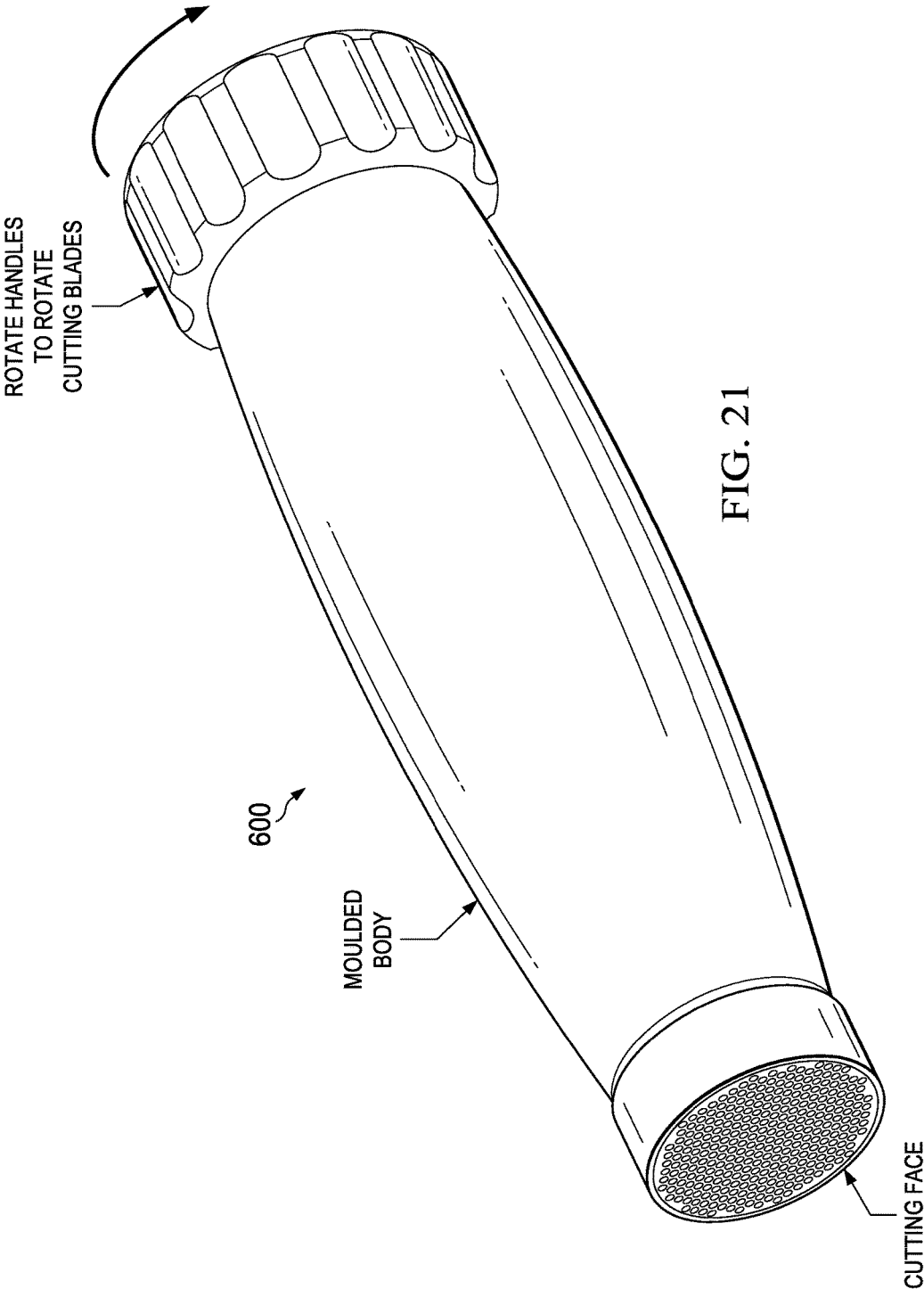


FIG. 20



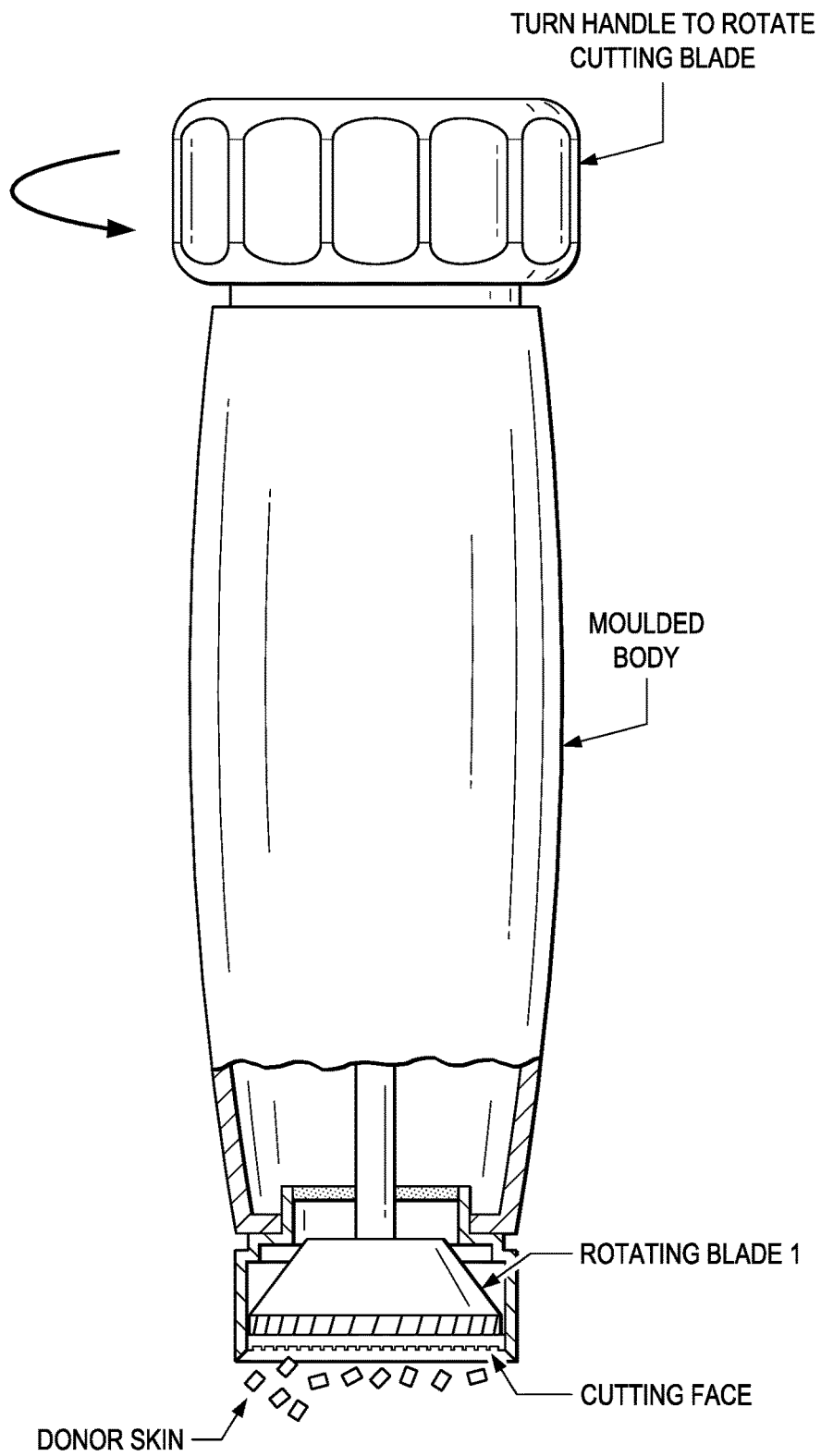


FIG. 22

## SKIN GRAFT DEVICES AND METHODS

## RELATED APPLICATIONS

This application is a continuation of U.S. patent application Ser. No. 13/451,174, filed Apr. 19, 2012, which claims the benefit under 35 USC 119(e), of the filing of U.S. Provisional Patent Application No. 61/477,485, entitled "Skin Graft Devices and Methods," filed Apr. 20, 2011, the entire contents of which are incorporated herein by reference.

## BACKGROUND

## 1. Field of the Invention

The present invention relates generally to skin grafting and related devices and methods. The present invention provides a systematic approach to the process of skin grafting, i.e., harvesting, post-excision processing and application of donor skin and treatment of the graft recipient site.

## 2. Background Information

Advances in medical technology have provided many patients with benefits inconceivable a century ago. In particular, skin grafting has enabled doctors to heal wounds with the patient's own skin from a harvest site on the patient. The skin grafting techniques have many wonderful benefits, but are still replete with a number of problems.

The process of split-thickness skin grafting can be envisaged as a series of steps; (1) harvesting the split-thickness-skin graft ("STSG") at a donor site; (2) processing of excised STSG; (3) application of the processed skin to the wound site; and (4) pre- and/or post-graft treatment to accelerate healing of the wound site. Each of these steps interposes various challenges and obstacles, e.g., technical, therapeutic and financial, in executing a successful graft.

In regard to the first step, harvesting a STSG at a donor site has traditionally been accomplished using powered, hand-held dermatomes. These devices are expensive and the operation is known to be highly dependent on user skill and training, and requires involved procedures to accurately obtain a successful harvest. These devices must be operated at a precise constant angle relative to the skin, with the exact amount of pressure to insure a uniform harvest. Slight variations in operative use of these dermatomes result in excised skin of variable-thickness, which sometimes must be discarded altogether. As a result, these devices are primarily wielded only by experienced plastic surgeons. Use of these dermatomes are generally confined to the operating room setting, increasing the cost of the procedure, especially given the average fee for operating room use.

There is a current need for harvesting procedures that require a lower degree of operator skill and are capable of being performed outside of an operating room, thus decreasing the costs of the procedure.

In regard to the second step of processing excised skin, it is highly desirable to maximize the coverage of the donor skin at the wound site for any given area of a potential donor site. Apart from minimizing trauma incurred at the donor site, a major factor limiting survival following extensive injury is insufficient availability of donor sites to provide enough skin for the required grafting procedures. One procedure is to mesh the skin graft i.e., creating slits in the excised donor skin to allow for the skin to be stretched. A graft-meshing machine is commonly used in hospital-based surgical practices, and generally allow for an expansion ratio of 3:1 to 9:1. The excised harvested skin is placed on a specific template, depending on the expansion ratio desired,

and the template and graft are pressed through the mesher. While greater ratios than 9:1 may be possible using meshing techniques, there is a concomitant significant delay in epithelialization with using such ratios. When healed, a meshed grafted site characteristically has a permanent "crocodile skin" or "weaved" appearance.

Micro grafting techniques, in which the donor tissue is actually minced in order to achieve a greater than 10:1 expansion ratio, are known in the art. Such techniques allow for a greater coverage area from a small donor site than meshing techniques. Traditional micrograft techniques, dating back to 1963, utilized minced skin that is between 1/8th inch (approximately 3 mm, or 3000  $\mu$ m) and 1/16th inch (approximately 1.5 mm, or 1500  $\mu$ m) in size. However, disadvantages of using pieces larger than 1500  $\mu$ m have been noted. For example, in skin pieces of this size cells remote from a cut edge have a limited availability to migrate and proliferate and thereby contribute to forming new skin. In addition, the techniques employed have required each piece to be oriented epidermis upwards, making the procedure tedious and impractical. Further, the appearance of the new skin that is produced using particles of this size is poor, often having a cobblestone appearance.

There is currently a need for a procedure capable of producing micrograft particles in a size less than 1500  $\mu$ m in a rapid and efficient manner, with a minimum of handling procedures, while resulting in skin pieces that are viable and capable of "taking" when applied to a wound site. Such technique would significantly aid in the ease and speed of operations utilizing micrografts.

The third step of the graft procedure, application of processed excised skin to the wound site, it is particularly relevant to the application of micrograft particles to a wound site. Current methods of distributing micrografts, such as mechanical spreading results in clumps or aggregates of skin particles, frustrating an even distribution. In addition, in larger aggregates, some micrograft particles will not be in direct contact with the wound bed. Such particles cannot readily integrate with the wound bed and also will have a reduced potential for nourishment from the wound fluid exudates and consequently have a decreased potential to remain viable. Thus, the aggregation of micrografts reduces the efficiency of epithelialization and may significantly increase the time required to close a wound.

There is a current need for devices and methods to effect an even distribution of micrograft particles on a wound surface, thereby promoting the efficiency of epithelialization.

The fourth step of the graft procedure relates to pre- and/or post-graft treatment to accelerate healing of the wound site. As is known in the art, closure of surface wounds involves the inward migration of epithelial, dermal and subcutaneous tissue adjacent to the wound. This migration is ordinarily assisted through the inflammatory process, whereby blood flow is increased and various functional cell types are activated. Through the inflammatory process, blood flow through damaged or broken vessels is stopped by capillary level occlusion; thereafter, cleanup and rebuilding operations may begin.

## SUMMARY

Certain embodiments of the present disclosure comprise devices and methods relating generally to skin grafting. Particular embodiments provide a systematic approach to

the process of skin grafting, i.e., harvesting, post-excision processing and application of donor skin and treatment of the graft recipient site.

Certain embodiments of the present disclosure comprise a device for obtaining a plurality of skin tissue particles for use in skin grafting. In particular embodiments, the device includes a dressing comprising a first surface configured to retain a plurality of skin tissue particles; a housing configured to receive the dressing, wherein the housing comprises a first aperture configured to be coupled to a vacuum source; and a plurality of hollow needles proximal to the first surface of the dressing. In certain embodiments, the first surface comprises a gel. In specific embodiments, the gel is a polyurethane film, an extra-cellular matrix (e.g. collagen), or a silicone based polymer.

In particular embodiments, the plurality of hollow needles are tapered. In specific embodiments, the plurality of hollow needles are tapered such that each of the plurality of needles comprises a larger end proximal to the first surface of the dressing. In certain embodiments, the housing comprises a seal configured to extend around the plurality of hollow needles. In particular embodiments, during use, the plurality of hollow needles are placed proximal to a donor site contained within the seal.

In specific embodiments, the dressing is located between the plurality of hollow needles and the first aperture. In particular embodiments, the dressing is removable from the housing, and in certain embodiments, the housing comprises a first aperture configured to be coupled to a bellows.

Exemplary embodiments also comprise a method of obtaining a plurality of skin tissue particles for use in skin grafting, the method comprising: placing a first device according to claim 1 onto a first donor site; applying negative pressure to the first device; removing a first plurality of skin tissue particles from the first donor site; removing a first dressing from the housing of the first device, wherein the first plurality of skin tissue particles are retained on the first surface of the dressing; and placing the first dressing on a graft site, where the first plurality of skin tissue particles are proximal to the graft site.

Particular embodiments may also comprise covering the dressing and the graft site with a drape; and applying negative pressure to a region under the drape. Certain embodiments may further comprise: placing a second device onto the donor site and applying negative pressure to the second device; removing a second plurality of skin tissue particles from the donor site; removing the second dressing from the housing of the second device, wherein the second plurality of skin tissue particles are retained on the first surface of the second dressing; and placing the second dressing on the graft site, wherein the second plurality of skin tissue particles are proximal to the graft site.

Certain embodiments may also comprise a device for obtaining a plurality of skin tissue particles for use in skin grafting, the device comprising: a processor configured to process skin tissue into a plurality of skin tissue particles; and a container configured to retain the plurality of skin tissue particles, wherein the processor comprises a first cutting surface configured to penetrate skin tissue at a donor site and a second cutting surface that rotates.

In particular embodiments, the first surface comprises a punch configured to penetrate skin tissue and the second cutting surface is configured to sever skin tissue from the donor site. In certain embodiments, the second cutting surface rotates in a plane generally parallel to the skin tissue of the donor site. In specific embodiments, the second cutting surface rotates in a plane generally perpendicular to

the skin tissue. In particular embodiments, the processor is manually operated. In certain embodiments, the processor is electrically powered. In specific embodiments, the device is configured such that the device can be separated from the processor.

The following drawings illustrate by way of example and not limitation. For the sake of brevity and clarity, every feature of a given structure is not always labeled in every figure in which that structure appears. Identical reference numbers do not necessarily indicate an identical structure. Rather, the same reference number may be used to indicate a similar feature or a feature with similar functionality, as may non-identical reference numbers.

#### BRIEF DESCRIPTION OF THE DRAWINGS

FIGS. 1-5 illustrate perspective and section views of a first exemplary embodiment.

FIGS. 6-14 illustrate section views of the embodiment of FIGS. 1-5 during use.

FIGS. 15-18 illustrate perspective and orthographic views of a second exemplary embodiment.

FIGS. 19-20 illustrate perspective and orthographic views of a third exemplary embodiment.

FIGS. 21-22 illustrate perspective and orthographic views of a fourth exemplary embodiment.

#### DESCRIPTION OF ILLUSTRATIVE EMBODIMENTS

The term “coupled” is defined as connected, although not necessarily directly, and not necessarily mechanically; two items that are “coupled” may be integral with each other. The terms “a” and “an” are defined as one or more unless this disclosure explicitly requires otherwise. The terms “substantially,” “approximately,” and “about” are defined as largely but not necessarily wholly what is specified, as understood by a person of ordinary skill in the art.

The terms “comprise” (and any form of comprise, such as “comprises” and “comprising”), “have” (and any form of have, such as “has” and “having”), “include” (and any form of include, such as “includes” and “including”) and “contain” (and any form of contain, such as “contains” and “containing”) are open-ended linking verbs. As a result, a wound-treatment method that “comprises,” “has,” “includes” or “contains” one or more steps possesses those one or more steps, but is not limited to possessing only those one or more steps. Likewise, a wound dressing that “comprises,” “has,” “includes” or “contains” one or more elements possesses those one or more elements, but is not limited to possessing only those elements.

Further, a device or structure that is configured in a certain way is configured in at least that way, but it can also be configured in other ways than those specifically described.

Referring now to FIGS. 1-5, a device 100 for obtaining skin tissue particles for use in skin grafting is shown. Device 100 comprises a housing 110 that is configured to receive a dressing 120. In certain embodiments, housing 110 may comprise a first portion 111 and a second portion 112 that may be coupled together or separated to allow the contents within housing 110 to be removed. In the embodiment shown, device 100 also comprises a plurality of hollow needles 130, which are configured to penetrate tissue during use and remove skin particles from a harvest site. In the illustrated embodiment, device 100 further comprises a seal 150 configured to seal an area around the plurality of needles 130.

An overview of the operation of device **100** will be provided initially, followed by a more detailed description. During operation, device **100** can be placed on a skin harvest site such that hollow needles **130** are in contact with the harvest site and seal **150** has sealed the area within the harvest site. A low pressure source **116** (e.g. a bellows device, a vacuum pump or other suitable device) can be coupled to aperture **140** and operated to provide a low pressure region within housing **110**. In the embodiment shown, a switch **115** can control operation of low pressure source **116**.

The operation of low pressure source **116** can cause air to flow in the direction of arrow **145** (see FIG. 3) and draw the harvest site toward needles **130**. Skin tissue from the harvest site can be drawn into needles **130** and removed from the harvest site. In certain embodiments, device **100** may be vibrated or laterally translated to assist in the removal skin tissue from the harvest site. In particular embodiments, hollow needles **130** may be tapered as shown in the cross-section view of FIG. 5 such that an end **131** proximal to dressing **120** is larger than an end **132** distal from dressing **120**.

When the skin tissue is removed from the harvest site, the tissue particles contact dressing **120**, which includes a surface **121** that is configured to retain the particles when dressing **120** is removed from housing **110**. A perspective view of dressing **120** removed from housing **110** is shown in FIG. 4. Dressing **120** can then be placed on a graft site oriented such that surface **121** and the tissue particles are in contact with the graft site. In certain embodiments, a negative pressure wound therapy can then be applied to the graft site to assist in the grafting of the tissue particles to the graft site.

Referring now to FIGS. 6-13, a description of individual steps in an exemplary method of use will be provided. As shown in FIG. 6, device **100** has been placed on a donor site **200** such that needles **130** are in contact with the skin tissue at donor site **200** and seal **150** has sealed the area of donor site around needles **130**. Referring now to FIG. 7, a negative pressure source (in fluid communication with device **100** via aperture **140**) has been operated to provide a negative pressure to device **100**. With negative pressure applied, tissue from donor site **200** is drawn into needles **130** to a controlled depth so that the tissue extends through needles **130** and to surface **121** of dressing **120**. Device **100** can then be manipulated (e.g. vibrated, laterally translated, or other suitable action) to remove skin tissue particles **131** from donor site **200**. When device **100** is removed from donor site **200**, skin tissue particles **131** are retained by surface **121** of dressing **120**. In particular embodiments, surface **121** may comprise a gel configured to retain the skin tissue particles by adhering the particles to the layer.

As shown in FIG. 8, device **100** has been removed from donor site **200** and surface **121** has retained skin tissue particles **131** after removal from donor site **200**. Referring now to FIG. 9, first portion **111** of device **100** has been separated from second portion **112**. This can allow dressing **120** to be removed from housing **110** of device **100**. As shown in this embodiment, surface **121** continues to retain skin tissue particles **131** with dressing **120** removed from housing **110**.

Referring now to FIG. 10, dressing **120** is positioned proximal to a graft site **300** so that skin tissue particles **131** are oriented proximal to graft site **300**. As shown in FIG. 11, in certain embodiments, multiple dressings **120** may be placed adjacent to each other in cases where graft site **300** is larger than dressing **120**. In specific embodiments, dress-

ing **120** may be configured so that multiple dressings can be placed adjacent to each other to substantially cover graft site **300**. For example, in certain embodiments, dressing **120** and surface **121** may be triangular, square, rectangular, hexagonal or other suitable configurations (when viewed from above looking down toward dressing **120** and graft site **300**).

As shown in FIG. 12, dressing **120** is placed on graft site **300** so that skin particles **131** are in contact with graft site **300**. As illustrated in FIGS. 8-12, skin tissue particles **131** are retained by surface **121** from the time the particles are harvested from donor site **200** until they are placed on graft site **300**. The ability to retain skin tissue particles **131** during the harvesting and grafting process provides many benefits, including the ability to provide a consistent and repeatable delivery of the particles to the graft site. This can increase the likelihood of obtaining a successful graft and improve the uniformity of the graft appearance.

In certain embodiments, negative pressure wound therapy can also be applied to graft site **300**. As shown in FIG. 13, a drape **160** has been applied over dressing **120** and graft site **300**. In the embodiment shown a coupling member **161** couples a conduit **162** to drape **160** and allows a negative pressure to be applied to the region under drape **160** (including e.g., dressing **120** and graft site **300**). As shown in FIG. 14, a larger drape **160** may be used in cases where multiple dressings **120** are needed to cover graft site **300**.

Referring now to FIG. 15-18, an exemplary embodiment comprises a unitary dermatome and mincer single use multi-bladed micro punch **400** with a plurality of nano-punches configured to incise the epidermis to the required depth and dimension. The punch can be activated using the snap switch situated on the side of the device. In this embodiment, a disposable tip also incorporates a specimen tray or container, where the processed skin particles are collected. Harvested material is released from the device by pressing the knob on the rear of the device. This embodiment provides a manual operation multi-specimen micro punch device with disposable tip and specimen tray/container. This device is able to harvest the skin particles (or islets) and process them to a ready to use state using simple operations. Additional details regarding the features and operation of the embodiment of FIG. 15-18 can be found in the figures and accompanying description in FIGS. 15-18.

Referring now to FIGS. 19-20, a unitary, battery powered, re-useable dermatome and mincer is shown. In this embodiment, the battery-powered device is placed on the donor site with moderate pressure and activated using the on/off trigger. During operation of this embodiment, the skin tissue can be excised and transferred by vacuum to a removable/disposable collection canister or container. In this exemplary embodiment, the cutting head may also be a disposable component. The canister/container of this device provide easy access to fully processed skin particles. Additional details regarding the features and operation of the embodiment of FIG. 19-20 can be found in the figures and accompanying description in FIGS. 19-20.

Referring now to FIGS. 21-22, a disposable, single-use skin mincer is illustrated. In this embodiment, the donor skin can be harvested using a dermatome or Weck knife and then placed into the device. Exemplary embodiments of this device comprise two contra-rotating blades which are operated by rotating the handle, which grind the donor skin tissue into small particles. The finished skin particles can be ejected from the device once the required dimensions have been achieved. This embodiment provides a unique method of processing the donor skin by using a twisting action to prepare the donor skin ready for grafting. Additional details



regarding the features and operation of the embodiment of FIG. 21-22 can be found in the figures and accompanying description in FIGS. 21-22.

The claims are not intended to include, and should not be interpreted to include, means-plus- or step-plus-function limitations, unless such a limitation is explicitly recited in a given claim using the phrase(s) “means for” or “step for,” respectively.

We claim:

1. A device for obtaining a plurality of skin tissue particles, the device comprising:
  - a first portion having a conical end and a cylindrical end coupled to a base of the conical end and an aperture in an apex of the conical end, the first portion having a flow path through the first portion from the aperture to the cylindrical end, the aperture configured to be coupled to a low pressure source;
  - a second portion having a first end configured to be coupled to the cylindrical end of the first portion to form a cavity, the second portion being cylindrical and having a second end that is closed, the second portion being separable from the first portion;
  - a plurality of needles coupled to the second end of the second portion, each of the plurality of needles being hollow and in fluid communication with the aperture when the second portion is coupled to the first portion; and

a dressing configured to be disposed in the cavity and in fluid communication with the aperture and the plurality of needles.

2. The device of claim 1, wherein each of the plurality of needles is tapered, an end of each needle proximate to the second end of the second portion being larger than a distal end.

3. The device of claim 1, further comprising a seal coupled to the second portion, the seal surrounding the plurality of needles.

4. The device of claim 1, wherein the first portion further comprises a surface extending across the cylindrical end, the surface having a plurality of apertures.

5. The device of claim 4, wherein the cavity is formed between the surface extending across the cylindrical end of the first portion and the second end of the second portion.

6. The device of claim 1, wherein the cylindrical end of the first portion has an outer diameter, the second portion has an outer diameter, and the outer diameter of the first portion is substantially the same as the outer diameter of the second portion.

7. The device of claim 6, wherein the second portion further comprises an annular tab having an outer diameter less than the outer diameter of the second portion to form an annular shoulder, the annular shoulder configured to receive the cylindrical end of the first portion.

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