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PHILIPPINE PATENT (19)

[11] No.:

[45] Issued SEP 28 1992

[54] Title: METHOD FOR DOUBLE DIPPING GELATIN COATED CAFLETS

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[22] Filed: December 2, 1988

[21] Application Serial No: 37825

FOREIGN APPLICATION PRIORITY DATA

[31] Number(s) : 129,109

[32] Date(s) : December 4, 1987

[33] Country(ies) : U. S. A.

[52] PH Class ..... 424/451; 424/37; 424/486; 514/774;  
428/403

[51] Int. Class ..... A61K 9/40;

[53] Field of Search ..... 424/451; 424/37; 424/486; 514/774;  
428/403  
A61K 9/40; B32B 9/02

[56] Reference(s) Cited and/or Considered: None

[57]

(see abstract next page)

BAD ORIGINAL



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METHOD FOR DOUBLE DIPPING GELATIN COATED CAPLETS

A b s t r a c t

5 A novel capsule-like medicament and method for producing such medicaments are disclosed. The method provides a procedure for coating solid cores, such as caplets, with a first gelatinous coating on one end, and then with a second gelatinous coating on the other end which is thicker than the first, to simulate the interlocking halves of a hollow capsule. The second, thicker gelatinous coating can be provided with a single gelatin coating from a bath having a higher viscosity than the bath used to provide the first gelatinous coating.

10 Alternatively, the second gelatinous coating can be provided by double dipping to provide layers of gelatinous material or gelatin.

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METHOD FOR DOUBLE DIPPING GELATIN COATED CAPLETS

CROSS-REFERENCE TO RELATED APPLICATIONS

5 This application is a continuation-in-part application of my co-pending application, Serial Number 016,914, filed February 20, 1987, entitled "GELATIN COATED CAPLETS AND PROCESS FOR MAKING SAME", which is commonly assigned and which is also hereby incorporated by reference.

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Field of the Invention

This invention relates to coated medicaments and processes for providing gelatinous coverings for such medicaments.

15 This invention is also directed to novel apparatus for producing such medicaments.

Background of the Invention

20 Until recently, the pharmaceutical industry has relied upon empty gelatin capsules for the encapsulation of medicinal agents as a popular method for administering drugs. Hard capsules are not new. As early as 1848 Murdock introduced the two-piece, hard gelatin capsule.

25 Capsules are tasteless, easily administered and easily filled either extemporaneously or in large quantities commercially. Many patients find it easier to swallow capsules than tablets, therefore preferring to take this form whenever possible. This preference has prompted

30 pharmaceutical manufacturers to market certain products in capsule form even though they are also available in tablet form.

Empty gelatin capsules are typically made from gelatin-glycerin, pure gelatin, starch or sugar gelatin,

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or other soluble gelatin combinations. See Remington's Practice of Pharmacy, Martin & Cook, 17th edition, pp. 1625-1630, which is herein incorporated by reference. Capsules serve as adequate housings for powders, masses, 5 liquids, pellets and oils and offer improve palatability and convenience.

Generally, empty hard gelatin capsules are manufactured using automated equipment. This equipment employs rows of 10 stainless steel pins, mounted on bars or plates, which are dipped into a gelatin solution maintained at a uniform temperature and fluidity. The pins are then withdrawn from the gelatin solution, are rotated and then inserted into drying kilns through which a strong blast of filtered 15 air with controlled humidity is forced. A crude capsule half is thus formed over each pin during drying. Each capsule half is then stripped, trimmed to uniform length, filled and joined to an appropriate mating half. Such hard capsule making systems are sold by Cherry-Burrell of 20 Cedar Rapids, IA.

During most of this century, empty gelatin capsules were a popular dosage form for prescription and over-the-counter (OTC) drugs. However, in the early 1980's there was an 25 unexpected increase in tampering with the contents of those capsules, resulting in several widely publicized deaths. This curtailed consumer demand for these products, caused ubiquitous concern regarding safety among those in the pharmaceutical community, and idled much of 30 the industry's hard capsule making equipment. Improved gelatin capsules and tamper-resistant packaging were then developed, but were expensive to produce and were not foolproof.

Once the threat of capsule tampering was recognized, many manufacturers withdrew their capsule products from the market, often replacing them with solid, oblong-shaped medicaments referred to commonly as caplets. Caplets are 5 solid oblong tablets which are sometimes coated with material such as cellulose. Typically, this coating is applied using coating-pan systems such as the "Vector-Freund Hi-Coaters, sold by Vector Corporation, 675 44th Street, Marion, Iowa, or the GC-1000" sold by Glatt 10 Air Techniques' 20 Spear Road, Ramsey, New Jersey.

A coating-pan system has a perforated pan or a drum which revolves in a manner similar to a standard clothes dryer. The system includes an air-atomization, spray gun which is 15 inserted into the center of the drum for spraying a fine mist of coating material. A batch of solid medicaments or caplets is typically introduced into the cylindrical pan, wherein said batch is caused to tumble. The tumbling action tends to smooth out some of the rough edges on the caplets prior to coating them with organic or aqueous film 20 solutions which may contain solid additives. Coating pans generally produce consistent coating thicknesses and weights but are capable of providing only one color 25 coating. Coatings produced by this method are often thin, offering poor coverage of medicament imperfections and rough edges not removed by the tumbling operation. Unless time is taken to build up a thicker coat, defects on the 30 solid core result in a medicament that does not exhibit a pleasing appearance and may be perceived as being harder to swallow. Moreover, coating abrasion occurring during tumbling produces a surface finish on these medicaments that fails to exhibit the shiny surface that consumers and those in the art have associated with ease of 35 swallowability. Applicant has pan coated caplets with gelatin on an experimental basis and has measured coating

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thicknesses of only about 6 mils. Moreover, these pan coated gelatin caplets were not observed to be as shiny as caplets coated by a dipping process.

5     Swallowability, the ability to pass through the fauces, pharynx and esophagus into the stomach, is dependent on the physical characteristics of the medicament as well as psychological factors. See Stedman's Medical Dictionary, Anderson Publishing Co., 5th edition' p.1377, which page  
10    is herein incorporated by reference. Physical characteristics, such as medicament shape, size and surface finish, can be correlated with esophageal adherence and swallowability. With respect to psychological factors, swallowing is normally volitional  
15    in adults, and muscular contractions of the throat are understood to be under the control of the individual at a subconscious level. See Stedman's, at 1377. Consumer surveys suggest that a shiny, capsule-like appearance has a special appeal to users as being easier to swallow. In  
20    addition, surveys indicate that consumers perceive capsule products to be more effective, thereby adding a possible additional placebo factor to their actual effectiveness.

Solid medicaments comprising gelatin in their coatings have been taught in a number of patents and abstracts.  
25    J.A. Glassman, U.S. Pat. No. 3,228,789, for instance, is directed to peroral capsules and tablets and a method for manufacturing same. Glassman discloses delayed release, compartmental medicaments with gelatin coatings, and  
30    includes treatments for tablets or pellet coatings. The abstract of Japanese patent 52041213, assigned to Feund Industry Ltd., discloses a process for coating tablets with a solution containing gelatin as a film-forming agent. The abstract of Japanese patent 69027916, assigned  
35    to Sankyo Co. Ltd., is directed to gelatin coated tablets

and a process for making same. The process of this patent includes feeding raw tablets at continuous intervals into a support. The tablets are immersed in a coating solution which can comprise gelatin. They are then recovered and held on a holder. Excess coating solution deposited at the lower surface of the tablet is removed by an eliminating plate, and finally the tablet is released into a cooling solution from which it is recovered and dried to produce a seamless coated tablet. The abstract of Japanese patent 65009992, assigned to Konishi, is directed to a film-coating method using gelatin for coating tablets in a coating pan. The gelatin described in this abstract is pre-treated with water in a pressure-cooker at a 120°-140°F for 30-40 minutes to reduce the adhesive properties of the gelatin to allow coating of the tablet. The abstract of Japanese patent 65009994, also assigned to Konishi, is directed to coating tablets in a coating pan with an emulsion including a mixture of hot water, gelatin, a surface active agent and a member selected from a group consisting of fats and oils, paraffin and wax. The use of the emulsion described in this patent abstract allows tablets to be coated with gelatin in the same manner as coating tablets with sugar. See also the abstract of an article by Richardson entitled "Franciscus Pill Coater", Phar. Hist., 28: 90-91 (2) 1986. This abstract is directed to the Franciscus Pill Coater, one of the later refinements of the gelatin-coated process that appealed to the practicing pharmacists in the 19th century. Other abstracts also disclosing coatings for solid medicaments comprising gelatin include those for: Japanese patent 60084215, assigned to Shinetsu Chemical Industries, U.S. Patent 4,238,510, assigned to Lifesavers, Inc., and Japanese patent 69026677, assigned to Daiichi Seiyaku Co. Ltd.

Several patents have disclosed the concept of coating pills by dipping half the surface of the pill at a time. Richards, U.S. Patent No. 599,865 is directed to a process and apparatus for dipping pills wherein an adhesive bearing bar is used to hold the pills before dipping them into gelatin. This process requires great care in maintaining the consistency of the adhesive material, i.e. wax, so that each pill will adhere to the dipping-bar. The specification of Richards also warns that great care must be taken not to dip the pills so deep as to get any of the gelatin upon the wax, which may ruin its adhesive capacity. The method of Richards additionally is labor intensive, and therefore, is more expensive by today's standards. Clark, U.S. Patent No. 724,436, is directed to a pill coating machine that employs pill-bars having a series of perforations for receiving pills. Each perforation is adapted for suction, whereby the pill is held in position during the dipping operation. Banker, U.S. Patent No. 3,896,762, discloses a rotary immersion coating that similarly employs suction to hold solid medicaments prior to passing these medicaments through a coating bath. While Clark and Banker provide apparatus for holding and dipping medicaments, neither discloses that the final product will exhibit a capsule-like appearance with or without a seam. Moreover, applicant has tested vacuum holding apparatus and has discovered that the suction tends to attract some of the gelatin into the holder, producing an irregular seam. Vacuum holding systems such as these also require significant power consumption, are often complicated and uncertain in their action, and necessitate expensive and sensitive vacuum equipment. Finally Oddo, U.S. Patent 3,045,641, discloses apparatus for color-coding tablets that utilizes a rotating resilient roller impregnated with a coating substance, whereby tablets are passed beneath the roller

on a conveyor and are deeply impressed into the resilient roller surface. This patent does not disclose the use of gelatin or the use of a dipping process to produce a thick capsule-like coating.

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Although these gelatin coated medicaments and processes have achieved some commercial success in the marketplace, a need remains for a coated medicament which is at least as tamper-resistant as a caplet while providing the ease of swallowability of a capsule. There is also a need for 10 a less expensive medicament coating method capable of producing a multi-colored, capsule-like coating which is perceived by the consuming public to be more effective.

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#### SUMMARY OF THE INVENTION

The present invention provides a novel method for coating solid cores such, as caplets, with gelatinous coatings to produce simulated capsule-medicaments. The gelatin coated medicaments are coated by individually dipping and drawing 20 first one end of the medicament, and then applying a more viscous or double gelatinous coating to the other end of the medicament to produce a capsule-like seam. Preferably, the thickness of the coating applied to the 25 first end of the medicament should be roughly twice as thick as the coating applied to the other end, to give the appearance and feel of a two-piece capsule having overlapping halves. The more capsule-like appearance has a special appeal to users as being easier to swallow. 30 Since the capsule-like seam of medicaments produced by the methods of this invention could be perceived by consumers as having the effectiveness of capsule products, there may be an additional placebo factor to their overall effectiveness. In the event that a fully tamper-proof 35 capsule cannot be developed in the near future, gelatin

coated caplets produced by the methods of this invention could partially satisfy the enormous consumer demand created by the withdrawal of many capsule products from the market place.

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Accordingly, a medicament is provided that is at least as tamper-resistant as a caplet while providing the ease of swallowability of a capsule. The various thicknesses of gelatinous coatings of this invention provide the appearance and feel of a capsule-like medicament.

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It is, therefore, an object of this invention to provide a simulated, capsule-like medicament having a gelatinous coating capable of being provided in two or more thicknesses.

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It is another object of this invention to provide a simulated, capsule-like medicament that is tamper resistant.

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It is another object of this invention to provide a simulated, capsule-like medicament that provides greater ease in swallowing and is perceived to be more effective than pan-coated pharmaceutical equivalents.

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It is still another object of this invention to provide a heavy layer of gelatin to cover imperfections inherent on the caplet core.

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With these and other objects in view, which will become apparent to one skilled in the art as the description proceeds, this invention resides in the novel construction, combination, arrangement of parts and

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methods substantially as hereinafter described and more particularly defined by the attached claims.

BRIEF DESCRIPTION OF THE DRAWINGS

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The accompanying drawings illustrate a complete embodiment of the invention according to the best mode so far devised for the practical application of the principles thereof, and in which:

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FIG. 1 is a diagrammatic view of the a preferred manufacturing sequence for providing a gelatin coating on caplets illustrating how the caplets are inserted, how the gelatinous coating is applied to the first and second ends 15 of the caplet, and how the caplets are dried and ejected;

FIG. 1A is a partial diagrammatic view of an alternative manufacturing sequence illustrating an alternative transferring method;

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FIG. 1B is a partial diagrammatic view of an alternative manufacturing sequence illustrating an alternative holding means and transferring method;

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FIG. 2 is an enlarged detail of a cross-sectional view of the holding means 26 of Fig. 1 with a caplet being dipped in a gelatinous material, said caplet being retained by "O"-rings 100 and 102;

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FIG. 3 is an enlarged detail of a transverse cross-sectional view of an alternate holding means illustrating a flat spring 202;

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FIG. 4 is an enlarged detail of a longitudinal cross-sectional view of a FIG. 3, taken through line 4-4, illustrating a caplet being retained by flat spring 202;

5 FIG. 5 is an enlarged detail of a transverse cross-sectional view of an alternative holding means embodiment illustrating a spring retention means 200;

FIG. 6 is an enlarged top view of an uncoated caplet;

10 FIG. 7 is an enlarged detail of a transverse view of the caplet of FIG. 6;

15 FIG. 8a-d are enlarged details of transverse views of caplets illustrating various coating patterns;

FIG. 9 is an enlarged longitudinal view of an alternative shape for an uncoated caplet;

20 FIG. 10 is an enlarged detail of a transverse view of the caplet of FIG. 9;

25 FIG. 11(a) and (b) are enlarged details of the longitudinal cross-section views of an alternative holding means embodiment showing how the ends of the caplet are held;

30 FIG. 12 is a partial diagrammatic view of an alternative manufacturing sequence whereby one end of the caplet is provided with a second gelatin coating; and

FIG. 13 is a partial diagrammatic view of an alternative manufacturing sequence illustrating how a second

gelatinous coating is applied to one end of the caplet without an intermediate kiln-drying step.

DETAILED DESCRIPTION OF THE INVENTION

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The present invention provides novel methods for coating caplets with gelatinous coatings to produce simulated capsule-like medicaments. The subject methods may be performed by modifying existing machines originally 10 intended to fabricate empty gelatin capsules or by newer similarly designed apparatus.

The novel processes of this invention comprise the steps of providing a holding means having a caplet channel 15 defined therein and inserting a first end of a caplet into said caplet channel while leaving the second end of the caplet exposed. The holding means is then manipulated relative to a bath of gelatinous material to dip the second exposed end of each caplet into that bath. The 20 resulting gelatinous coating on the second exposed end of the caplet is then permitted, and preferably caused, to dry to form a coated end. During the drying process, the caplet may be rotated to assist in uniformly distributing gelatin during drying. Once dry, the coated (second) end 25 of the caplet is then displaced through the caplet channel to expose its uncoated first end. A gelatinous coating is then applied to the uncoated first end of said caplet. The coating applied to the first end of the caplet is then permitted (or preferably caused) to dry, again with 30 rotation if desired for the purpose of spreading the coating evenly. In accordance with the preferred methods, the baths of gelatinous material into which the caplet ends are dipped, may be of different colors, to thereby create a simulated 2-piece capsule look to the finished 35 caplets with seams along their transverse axes. In a

preferred embodiment, one end of the caplet is double-dipped, or dipped in a more viscous gelatinous material, to create the appearance of overlapping capsule halves.

5 A substantial advantage of the present invention is that existing hard capsule manufacturing equipment may be readily adapted for the purpose of producing the coated caplet products of the present invention. In the  
10 preferred apparatus of the present invention, the conventional bars of such machines, having stainless steel capsule-forming protuberances mounted thereon, are replaced with bars having a plurality of cylindrical holding means mounted thereon. Each holding means  
15 receives, retains and facilitates the transfer of an individual caplet. The apparatus is fitted with a caplet feeder to feed caplets into each holding means. The holding means may, for example, be a cylinder which is open at both ends and which comprises a retaining means,  
20 such as "O"-rings or a spring biased retainer for the purpose of holding each caplet in position during the dipping process. The feeding means is preferably associated with an inserting means, which may be a simple channel and plunger assembly, for inserting a first end of  
25 each caplet into an appropriate holding means. The feeding means ensures that each caplet is inserted a sufficient distance to cause the second end of the caplet to appropriately protrude therefrom during the upcoming dipping process. Once each bar is loaded with caplets, it  
30 then proceeds to a dip station where the gelatinous coating is applied to the exposed ends of the caplets protruding therefrom, whereupon the bar is rotated through a first drying means for permitting the gelatinous coating to dry to form a coated second end. In a preferred apparatus, the second gripping means also comprises  
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substantially cylindrical holders which are open at both ends, and which have central bores defined therethrough. In this embodiment, these second holders are axially aligned with the bores of the first holders, at the 5 transfer positions, whereupon a plunger or other means is used to displace the half-coated caplets through and out of the "backs" of the first holds and into the "backs" of the second holders, and then through the second holders until the remaining uncoated ends of the caplets are 10 exposed for subsequent dipping. The dipping and drying processes are then repeated (preferably with a different colored gelatinous coating), whereupon a caplet ejection means pushes the caplets out of the second holders.

15 In another preferred embodiment of this method, the "fronts" of the second holder means are aligned with the "fronts" of the first holder means, whereupon the caplets are mechanically transferred from the first to the second holders without the need for an additional alignment 20 device. In still another embodiment, a single holding means is used for dipping both ends of the caplet, whereby, after dipping the second end, the caplet is transferred through this single holding means to expose the uncoated first end. This holder is then shifted to 25 the second gelatinous coating bath which preferably contains a different color gelatin for dipping the first end of the caplet.

The subject apparatus and method may be further understood 30 with reference to the figures. FIG. 1 illustrates, diagrammatically, the preferred method and apparatus for dipping solid caplets into a gelatinous material embodying the teachings of this invention. A holding means is first provided having a caplet channel 106 defined therein, 35 which can take any form having a cross-section sized to

slidably mate with said caplet 16. The caplet 16 having a first and second end, 110 and 104 of FIG 2, is inserted using inserting means 20 into said caplet channel 106 while leaving the second end of the caplet 104 exposed.

5 Next, a gelatinous coating, known to those in the pharmaceutical arts, is applied by first application means 28 to the exposed second end 104 of the caplet. The depth to which the second end 104 can be coated is dependant upon the desired color configuration and seam requirements. The half-coated caplet is then dried using the first drying means 30 and 32 which permits the gelatinous coating on the second end 104 to dry, forming a coated second end. The caplet 16 is then displaced through said caplet channel 106 to expose the first end 110 preferably using a gripping means illustrated in the embodiment of FIG. 1 as transfer means 12, alignment means 18 and second holding means 15. The caplet 16 is then coated with a gelatinous material on its first end 110 by second application means 38 which is then dried by second drying means 34 and 36, resulting in a dry caplet substantially covered in gelatin. Accordingly, this invention provides novel means for providing simple and inexpensive modifications to existing hard capsule equipment to manufacture simulated capsule-like medicaments. This invention teaches preferred process sequences that supplement and partially replace otherwise standard empty gelatin capsule techniques and parameters that are known to the art. For example, gelatin materials used for the coatings of this invention may be any of the well known types utilized in the art of manufacturing empty capsules and coated medicaments.

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Referring again to FIG. 1, the caplet 16 is fed into inserting means 20 by feeder 11. The feeder 11 preferably comprises a chute attached to a reservoir. Alternatively,

mechanical means or pneumatic means may be developed for this purpose. In one embodiment, a 20-40 wide channel vibrational feeder has been deemed useful. It is expected that those in the art could readily adapt current automation technology to develop a means for feeding caplets into inserting means 20 of this invention. In the preferred embodiment, plungers displace caplets into each of a series of molding means spaced apart along a bar mount, shown in end cross section in FIG. 1, i.e., holding means 26. Each caplet 16 is then inserted into a first holding means 26 using plunger 20. Both the first and second holding means 26 and 15 are preferably cylindrical, having caplet channels 106 having a cross-section sized to slidably mate with the caplet 16 to permit passage of the caplet 16. However, in the embodiment of FIG. 1A, these channels, or more preferably, bores, can extend through the holding means with one cross-section sized to slidably mate with caplet and another cross-section sized to receive only a plunging means. This design is made possible due to the fact that the caplets can be transferred without displacing them through the entire length of the holding means in the method embodiment of FIG. 1A.

Included with one holding means embodiment of this invention, are retaining means for retaining the caplet at least during the first gelatinous coating application step. As shown by the embodiments found in FIGS. 2-5, the retaining means can comprise a plurality of recessed rubber "O"-rings 100 and 102, a flat spring 202 fixed by pin 204 and having convex side facing the caplet, or a resilient spring 200. As shown in FIG. 3 the flat spring 202 may be extended to enable external manipulation of the retaining means. The choice of retaining means is not critical and any known securing or resilient device may be

used. However, it is important that the retaining means provide enough clearance to pass the gelatin coated end, yet securely hold the uncoated end.

- 5 Inserting means 20 preferably comprises a plunging means having at least an end portion 22 disposed to abut the caplet 16 to effect displacement of the caplet 16 into the bore 106 of the first holding means 26.
- 10 In a preferred embodiment, a plurality of holding means are mounted on a fixture which can be transferred by the mechanical pushing means of a hard gelatin capsule assembly line. In such an embodiment, inserting means 20 has multiple plunging means having a plurality of end portions 22 disposed to abut multiple caplets to effect displacement thereof into the fixture. In one embodiment, 10 to 50 holding means, preferably 20 to 40, and most preferably 30 holding means are attached to the fixture. A plurality of said fixtures can be fed into a
- 15 20 conventional hard-capsule manufacturing assembly which can accommodate about 1500 to 1800 fixtures at a time.

The caplet coating process of FIG. 1 next applies a gelatinous coating to the second exposed end 104 of said caplet 16. A first application means 28 is employed for this purpose. In the preferred embodiment of this invention, groups of 4 or more fixtures are fed into a dipping means and vertically lowered into a gelatinous material such as methyl cellulose, calcium alginate or gelatin. The depth of the dip is preferably cam-regulated to the desired capsule size, color scheme, and "seam" requirements. As indicated in FIG. 8a-d, the color scheme can be bifurcated as depicted by caplet coatings 304 and 306, and a seam 302 or 300 can be provided by overlapping

the gelatinous coatings on the first and second ends 110 and 104.

5 The coatings on the first and second ends 110 and 104 of the caplet, when preferably dipped in gelatin can include plasticizer such as glycerin or sorbitol, water, preservatives, coloring agents, and opacifying agents.

10 See Remington's Practice of Pharmacy, pages 1625 to 1630. The preferred gelatin solution should be maintained at a uniform temperature and a constant degree of fluidity. If the gelatin solution varies in viscosity, it will correspondingly decrease or increase the thickness of the coating. Acceptable gelatin compositions can contain small amounts of methyl cellulose, polyvinyl alcohols, and 15 denatured gelatins to modify their solubility or produce a enteric effect. Common sources of gelatin contemplated by this invention include animal bones, hide portions and frozen pork skin. Grades of gelatin that are appropriate for this invention include pharmaceutical grade, food grade, Type A and Type B. Although the coatings herein provided can be made from any of these sources or grades, those learned in the art of capsule making are aware that the usual practice is to use a mixture of grades and 20 sources as dictated by availability and cost 25 considerations. Differences in the physical properties of finished capsules as a function of the type of gelatin used are slight. Reference may also be made to "The Theory and Practice of Industrial Pharmacy", by Lackman, Liberman and King (1970) pages 389-398, published by Lea and Febiger, Philadelphia, Pennsylvania, said pages being 30 hereby incorporated by reference. In a preferred embodiment of this invention, a gelatin mixture is prepared using 40% by weight bone (150 bloom), 20% by weight hyde (245 bloom) and 40% pork skin (270 bloom). 35 This mixture has a viscosity of 500cp as measured on a

Brookfield Chromatograph, at an operating temperature of 130°F.

Coloring can be added to the coatings to produce opaque or 5 transparent colors such as red, white, pink, green, reddish brown, blue, yellow and black. Colored medicaments are necessary to give a specialty product a distinctive appearance. Titanium dioxide is often added to the gelatin to form white medicaments, or to make an 10 opaque colored coating.

Still referring to FIG. 1, after coating the second end 104, the gelatinous coating is permitted to dry to form a coated second end. It is important to the teachings of 15 this invention that the caplet 16 is permitted to dry without contacting other objects, thus producing a shiny, simulated capsule-like finish on the caplet. In the preferred embodiment, a group of fixtures is raised from the gelatin and elevated to the first drying means, 20 comprising rotating means 30 and kiln means 32. Preferably, the caplets are rotated to distribute the coating on the caplet. In a most referred apparatus, the fixtures are automatically revolved after dipping to spread the gelatin more evenly over the caplet ends and 25 eliminate excess accrual at the ends. See Sindl, U.S. Patent No. 1,872,190, which is herein incorporated by reference. The caplets are then fed into a kiln drying means 32. Preferably, 5-60 fixtures containing caplets enter the drying kiln, where they move under drying 30 ducts. Air volume, temperature and humidity are controlled in the kiln and are set to conventional process parameters known to those in the industry.

When the gelatinous coating on the second end 104 of the 35 caplet is dry, it is displaced through the caplet channel

106 to expose the first end 110 using a gripping means illustrated in the embodiment of FIG. 1 as transfer means 12, alignment means 18 and second holding means 15. In one preferred embodiment, the transfer means 12 comprises 5 an end portion 14 disposed to abut said caplet to effect displacement of said caplet from the first holding means 26 to the second holding means 15. The caplet is then preferably displaced through the caplet channel or bore 106 of the first holding means 26, through the alignment means 18 and into said second holding means 15 to expose 10 the uncoated first end 110 of the caplet 16.

Alternatively, as depicted in FIG. 1A, when the gelatinous coating on the second end 104 is dry, it can be displaced 15 through the caplet channel 106 to expose the first end 110 by transferring the caplet from the "front" of a first holding means to the "front" of a second holding means, thus eliminating the need for alignment means 18 of FIG. 1. It is also envisioned that a single holding means 20 210 like the one illustrated in FIG. 11(a) and (b) could replace the use of the two holding means 26 and 15 of FIGS. 1 and 1A by providing for the displacement of the caplet through a central bore 201. Three "O"-rings 203, 205 and 207 are shown in FIGS. 11(a) and (b) for retaining 25 the caplet during the process of coating the second and first ends 104 and 110 using a single holding means 210. However, other retaining means, as previously described for the holding means 26 and 15 of FIG. 1, may also be employed for this purpose. As illustrated, in the 30 embodiment of FIG. 1B, by transferring bars containing a plurality of holding means 210 from the left side of the diagrammatic view of the process of FIG. 1 to the right side, the need for a second holding means is eliminated.

After displacing the caplet through the caplet channel 106, the protruding first end is ready for the application of a gelatinous coating which is illustrated on the right side of FIG. 1. As indicated in the above preferred 5 embodiments, groups of 4 or more fixtures can be fed into a dipping means 38 and vertically lowered into a gelatinous material, preferably containing a different color dye or pigment for providing a distinctive appearance. As indicated in FIG. 8, a seam 300 or 302 can 10 now be provided by overlapping the dried coating on the second end 104. Through careful selection of gelatin color schemes, the seam can exhibit a different color than the ends of the caplet, i.e., green and yellow coatings on the ends can be overlapped to form a blue seam. The 15 gelatinous coating on the first end is then permitted to dry without contacting other objects, as previously described for the second end 104. Separate rotating means 34 and kiln means 36 are illustrated in FIG. 1 for drying the coating on the first end. However, those in the art 20 may find it convenient to use the same drying apparatus used in drying the second end 104.

Finally, the caplet may be ejected from the second holder means 15 after the first end 110 is dry. Removal of the 25 coated caplet 16 can be effected by ejection means 25 which preferably is similar in structure to inserting means 20 and transfer means 12, in that it comprises an end portion 24 disposed to abut said caplet. Removing the caplet can be accomplished by plunging horizontally as in 30 FIGS. 1 and 1A or by plunging the caplet out of the holding means vertically as in FIG. 1B. The ejected caplet, now coated in gelatinous material, is then ready for printing and packaging.

In view of the above it is expected that a novel, simulated capsule-like medicament can be produced. The gelatin coated medicament of this invention which can be produced by the above method comprises a solid caplet 5 having a first and a second end, wherein a first gelatinous coating is provided on said second end, and a second gelatinous coating is provided on said first end of the caplet. The caplet generally is at least 2.5 times longer than it is wide, and ideally comprises a 10 cylindrical shape. The first and second gelatinous coatings substantially cover the caplet to form a simulated capsule-like medicament with a seam along a transverse axis of the medicament. As previously discussed, the first and second ends 110 and 104 of the 15 caplet can be coated with gelatinous coatings of different colors to provide a distinctive appearance for specialty products. A preferred color scheme for the medicament of this invention includes a caplet which is coated in a red and white gelatinous material. It has been discovered 20 that absorption of the gelatinous coating or the moisture in the gelatinous coating by the solid caplet may be reduced by applying a conventional precoat sealant to the caplet prior to dipping into a gelatinous material. See Baker, U.S. Patent 3,185,626, which is herein incorporated 25 by reference. Without a precoat sealant, it is possible that some of the gelatinous coating or moisture in the coating would seep into the caplet, resulting in a duller surface.

30 The gelatinous coatings of the previously described method are generally provided in substantially uniform thicknesses of about 5 to 40 mils, preferably about 10 to 30 mils, and most preferably from 15 to 25 mils. In accordance with an important aspect of this invention, a 35 procedure will now be described for double-dipping the

first end of the caplet to provide a combined thickness of about 10 to 80 mils, preferably about 20 to 60 mils, and most preferably about 30 to 50 mils. In accordance with this method, a holding means is provided having a caplet 5 channel defined therein. A first end of the caplet is then inserted into the caplet channel while leaving a second end of the caplet exposed. A gelatinous coating is then applied to the second exposed end of the caplet, and this coating is permitted to dry to form a coated second 10 end. Next, a first gelatinous coating is applied to the first end of the caplet whereby the gelatinous coatings on the first and second ends substantially cover the caplet. This first gelatinous coating on the first end of the caplet is then permitted to dry before a second gelatinous 15 coating is applied to the first end of the caplet. As used herein, the term "dry" need not require the absence of all moisture, and the caplet may even exhibit a wet appearance as long as a second gelatinous coating can be applied to the first end of the caplet without creating 20 significant visual imperfections in the coating. In accordance with this double dipping sequence, after the second gelatinous coating is applied, it is then permitted to dry so that the first and second gelatinous coatings on the first end form a thicker gelatinous coating than that 25 applied to the second end.

As in the previously disclosed single-dip procedure, the caplet may be pre-coated to reduce absorption of moisture and the gelatinous coatings by the caplet. Since 30 multi-colored medicaments are often preferred, the ends of the caplet can be provided with gelatinous coatings of different colors. The multiple coatings on the first end can be gelatinous coatings of similar colors, although this is not a requirement.

The double-dipping procedures described herein preferably include the same manufacturing sequence and apparatus as described previously for FIG. 1, with the exception that the caplet is not ejected until after a second gelatinous coating is applied to the first end of the caplet.

Referring now to FIG. 12, another double-dipping sequence is provided. After the caplet exits the kiln 36, it is provided with a second gelatinous coating on its first end by bath 139. Since the second gelatinous coating on the first end is preferably the same color and/or composition as the first gelatinous coating on the first end, it is anticipated that a single gelatinous bath can be used to provide both the first and second gelatinous coatings on the first end. In such an embodiment, the caplets would be returned to bath 38 after they exit the drying kiln 36. It is also envisioned that certain gelatinous materials could be applied successively without kiln-drying. As FIG. 13 illustrates, these compositions could be applied successively by merely applying a rotating step between the application steps.

After the second gelatinous coating is applied to the first end of the caplet, the caplet fixtures 34 are rotated to dry and/or distribute the coating on the caplet. Preferably a kiln drying apparatus 137 is employed to further dry the gelatinous materials after the rotating step. In accordance with both figures 12 and 13, the caplet can be ejected using plunger 125 after the gelatinous materials are sufficiently dry, leaving an empty holder 126.

In an important variation of this invention, a simulated capsule-like medicament is provided by providing a holding means having a caplet channel defined therein and

inserting a first end of the caplet into the caplet channel while leaving a second end of the caplet exposed. In further accordance with this method, a first gelatinous material is applied to the second exposed end of the 5 caplet to form a coated second end. The coated second end is permitted to dry prior to applying a second gelatinous material to the first end of the caplet. The second gelatinous material of this embodiment has a greater viscosity than the first gelatinous material so as to 10 provide a thicker gelatinous coating to said first end. Finally, the thicker gelatinous coating on the first end is permitted to dry. As in the case of the double-dipping sequence, a simulated capsule-like medicament is produced which is then preferably ejected using plunger 125.

15 The medicament provided in accordance with these embodiments will have a capsule-like seam generally along a transverse axis of the caplet and a length which is at least 2.5 times its width. This capsule-like medicament 20 also preferably uses gelatin as its gelatinous coatings. In the most preferred embodiment of the simulated capsule-like medicament of this invention, the second gelatinous coating partially overlaps the first gelatinous coating to form a capsule-like seam circumscribing the 25 medicament about a midway point of a longitudinal axis of the medicament.

Gelatin coated caplets can be supplied in a variety of shapes and sizes, from 000, the largest size which can be 30 swallowed, to 5, which is the smallest. Larger sizes can also be made available for use in veterinary medicine. FIGS. 6, 7, 9 and 10 illustrate two of the preferred shapes for caplets. FIGS. 6 and 7, show in top and transverse views respectively, an oblong caplet having a 35 raised portion circumscribing its perimeter. FIGS. 9 and

10 illustrate in longitudinal and transverse views another preferred embodiment having a cylindrical center portion and rounded ends having a transverse diameter slightly less than that of the cylindrical center portion. These 5 novel caplet designs facilitate the dipping method herein provided since they are easily held by the above retaining means and can be manufactured using conventional compression molding equipment.

10 From the foregoing it can be realized that this invention provides a simulated capsule-like medicament, a method for manufacturing this medicament, and apparatus used in the method. The advantages over the prior art are: increased tamper-resistance over hollow capsules, increased 15 swallowability over pan-coated medicaments, variable color scheme capability not available with pan-coated medicaments, less expensive operating costs and a greater perception by the consuming public that gelatin coated caplets are more effective. Although various embodiments 20 have been illustrated, this was for the purpose of describing, but not limiting, the invention. Various modifications, which will become apparent to one skilled in the art, are within the scope of this invention described in the attached claims.

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We claim:

1. A method for coating a caplet with a gelatinous coating to produce a simulated capsule-like medicament comprising:

5 (a) providing a holding means having a caplet channel defined therein;

(b) inserting a first end of said caplet into said caplet channel while leaving a second end of said caplet exposed;

10 (c) applying a gelatinous coating to said second exposed end of said caplet;

(d) permitting said gelatinous coating to dry form a coated second end;

15 (e) applying a first gelatinous coating to said first end of said caplet, said gelatinous coatings on said first and second ends substantially covering said caplet;

(f) permitting said first gelatinous coatings on said first end dry;

20 (g) applying a second gelatinous coating to said first end of said caplet; and

(i) permitting said second gelatinous coating on said first end to dry; said first and second gelatinous coatings on said first end forming a thicker gelatinous coating than that applied to said second end.

2. The method of claim 1 further comprising the step of providing a pre-coated caplet to reduce absorption of moisture and said gelatinous coatings by said caplet.

5 3. The method of claim 1 wherein said steps (c) and (f) coat said first and second ends of said caplet with gelatinous coatings of different colors.

10 4. The method of claim 1 wherein said steps (f) and (h) coat said first end in a gelatinous coating of a similar color.

15 5. The method of claim 1 wherein said steps (c) and (f) coat said caplet ends to a thickness of about 5 to 40 mils.

6. The method of claim 1 wherein said steps (c) and (f) coat said caplet ends to a thickness of about 10 to 30 mils.

20 7. The method of claim 1 wherein said steps (c) and (f) coat said caplet ends to a thickness of about 15 to 25 mils.

25 8. The method of claim 1 wherein said steps (f) and (h) coat said first end to a thickness of about 10 to 80 mils.

9. The method of claim 1 wherein said steps (f) and (h) coat said first end to a thickness of about 20 to 60 mils.

5 10. The method of claim 1 wherein said steps (f) and (h) coat said first end to a thickness of about 30 to 50 mils.

10 11. The method of claim 1, further comprising transferring said caplet from said first holding means to a second holding means.

15 12. The method of claim 11 wherein said caplet channel is selected to comprise a central bore extending through the length of said first holding means to permit passage of said caplet through at least a portion of said bore.

20 13. The method of claim 12 wherein said transferring step comprises displacing said caplet through said central bore.

25 14. The method of claim 13 wherein said transferring step comprises aligning said first and second holding means and displacing said caplet from said first holding means to said second holding means.

15. The method of claim 14 wherein said aligning step  
positions said second holding means proximate with said  
first holding means for displacing said caplet from the  
front of said first holding means to the front of said  
second holding means to expose said first end of said  
caplet.

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16. The method of claim 11 wherein said second holding  
means is selected to have a central bore extending through  
its length to permit passage of said caplet through at least  
a portion of said bore.

15  
17. The method of claim 16 wherein said transferring  
step displaces said caplet until said first end protrudes  
out of said second holding means.

18. The method of claim 1 wherein said gelatinous  
coating is selected to comprise gelatin.

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19. The method of claim 1 wherein said inserting step  
comprises disposing multiple caplets into a fixture comprising  
a plurality of holding means.

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20. The method of claim 19 wherein said inserting step  
comprises simultaneously introducing said multiple caplets  
into said fixture, said introducing step exposing a second  
end of said multiple caplets.

21. The method of claim 1 wherein said applying steps (c), (f) and (h) each comprise dipping said caplet into a gelatinous material and withdrawing said caplet, said dipping producing a wet, gelatinous coating on said first and second ends of said caplet.

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22. The method of claim 21 wherein said dipping step comprises vertically dipping said caplet into a bath of gelatinous material.

10

23. The method of claim 22 wherein said permitting steps (d), (g) and (i) comprise rotating said caplet to distribute said coating on said caplet.

15

24. The method of claim 22 wherein at least one of said permitting steps (d), (g) and (i) comprises kiln-drying said caplet.

20

25. The method of claim 11 further comprising the step of ejecting said caplet from said second holding means after said permitting step (i).

25

26. The method of claim 25 wherein said permitting step (g) comprises rotating said caplet to dry and distribute said first gelatinous coating on said first end.

27. The method of claim 26 wherein said permitting step (i) comprises rotating said caplet to dry and distribute said second gelatinous coating on said first end.

5 28. The method of claim 27 wherein said permitting step (i) further includes kiln-drying said caplet.

29. The method of claim 28 wherein said permitting step (g) further comprises kiln-drying said caplet.

10 30. The method of claim 28 wherein said applying step (f) and (h) comprise dipping said caplet into a single bath of gelatinous material.

15 31. The method of claim 28 wherein said applying steps (f) and (h) comprise dipping said caplet into different baths of gelatinous material.

20 32. The method of claim 29 wherein said applying steps (f) and (h) comprise dipping said caplet into a single bath of gelatinous material.

25 33. The method of claim 29 wherein said applying steps (f) and (h) comprise dipping said caplet into different baths of gelatinous material.

34. A simulated capsule-like medicament comprising:

(a) a solid caplet having a first and second end, said caplet comprising a generally cylindrical shape;

5 (b) a first gelatinous coating provide on said second end of said caplet; and

(c) a second gelatinous coating provided on said first end of said caplet, said second gelatinous coating having a greater thickness than the thickness of the said first gelatinous coating to form a simulated capsule-like medicament with a seam circumscribing the medicament about a 10 midway point of a longitudinal axis of the medicament.

15 35. The medicament of claim 34 wherein said second gelatinous coating comprises a plurality of gelatinous layers.

36. The medicament of claim 34 wherein said caplet is pre-coated to reduce absorbtion of moisture and said gelatinous coatings by said caplet.

20

37. The medicament of claim 34 wherein said first gelatinous coating is provided in a thickness of about 15 to 25 mils.

25

38. The medicament of claim 34 wherein said second gelatinous coating is provided in a thickness of about 30 to 50 mils.

39. The medicament of claim 34 wherein said first and second gelatinous coatings comprise gelatin.

40. The medicament of claim 35 wherein said plurality 5 of gelatinous layers have a combined thickness of about 30 to 50 mils.

41. The medicament of claim 37 wherein said first and second gelatinous coatings are provided in thicknesses of 10 about 5 to 40 mils.

42. The medicament of claim 37 wherein said caplet comprises a width and a length, wherein said length is at least 2.5 times the width.

15

43. The medicament of claim 37 wherein said first and second gelatinous coatings are provided in thicknesses of about 15 to 25 mils.

20

44. The medicament of claim 37 wherein said first and second gelatinous coatings comprise gelatin.

25

45. The medicament of claim 37 wherein said first second gelatinous partially overlaps said first gelatinous coating (partially overlaps said first gelatinous coating) forming a capsule-like seam circumscribing said medicament

-34-

at about a midway point of a longitudinal axis of said medicament.

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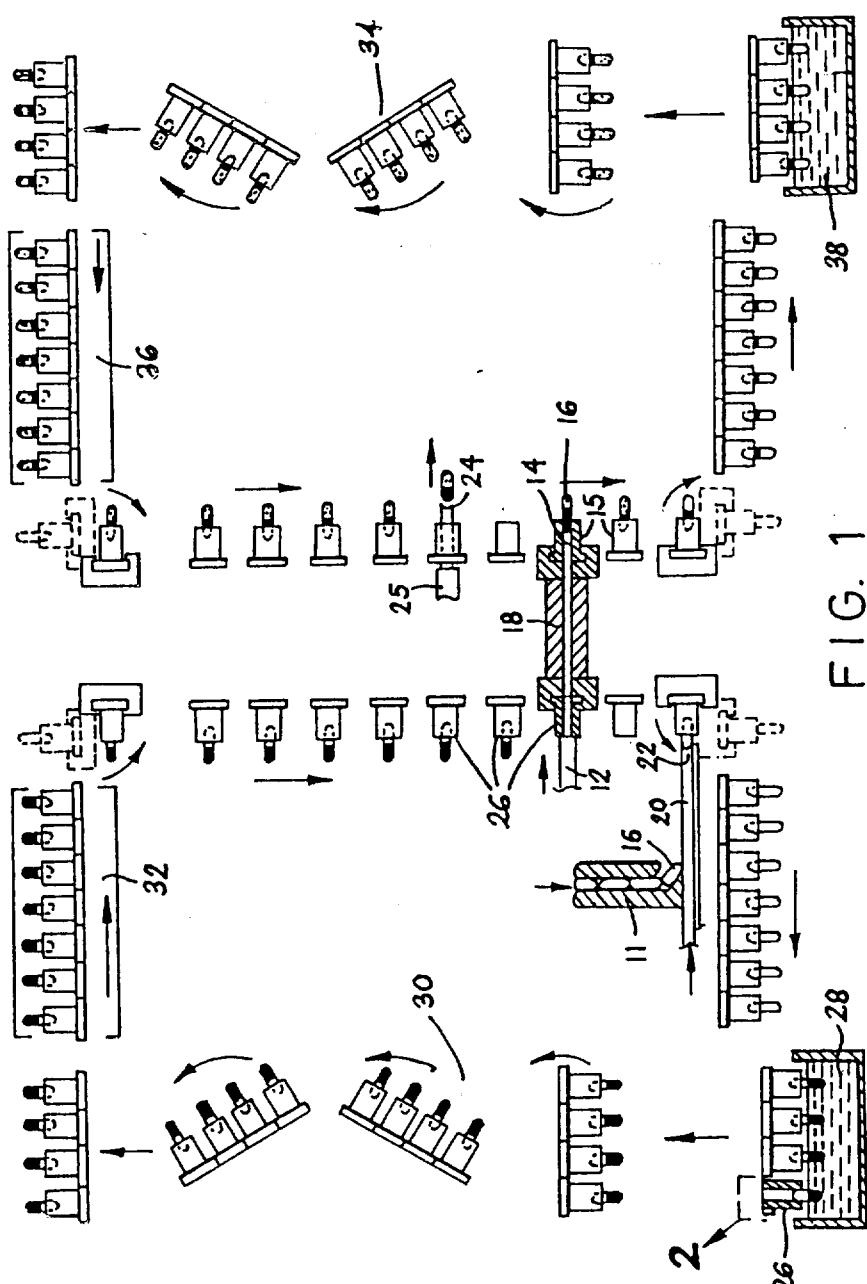


FIG. 1

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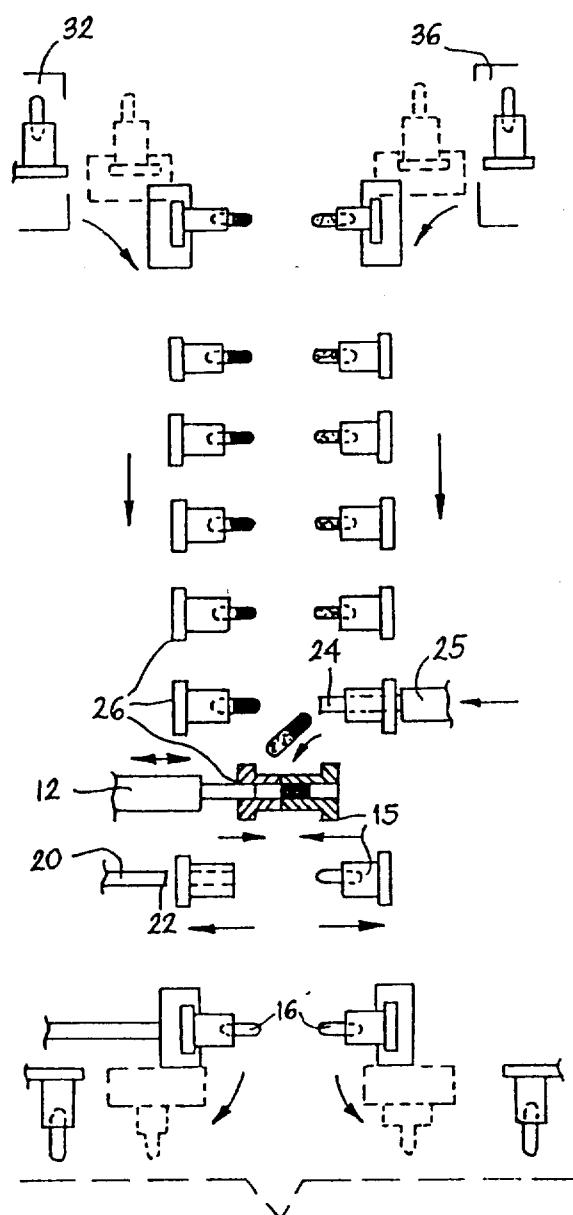


FIG. 1A

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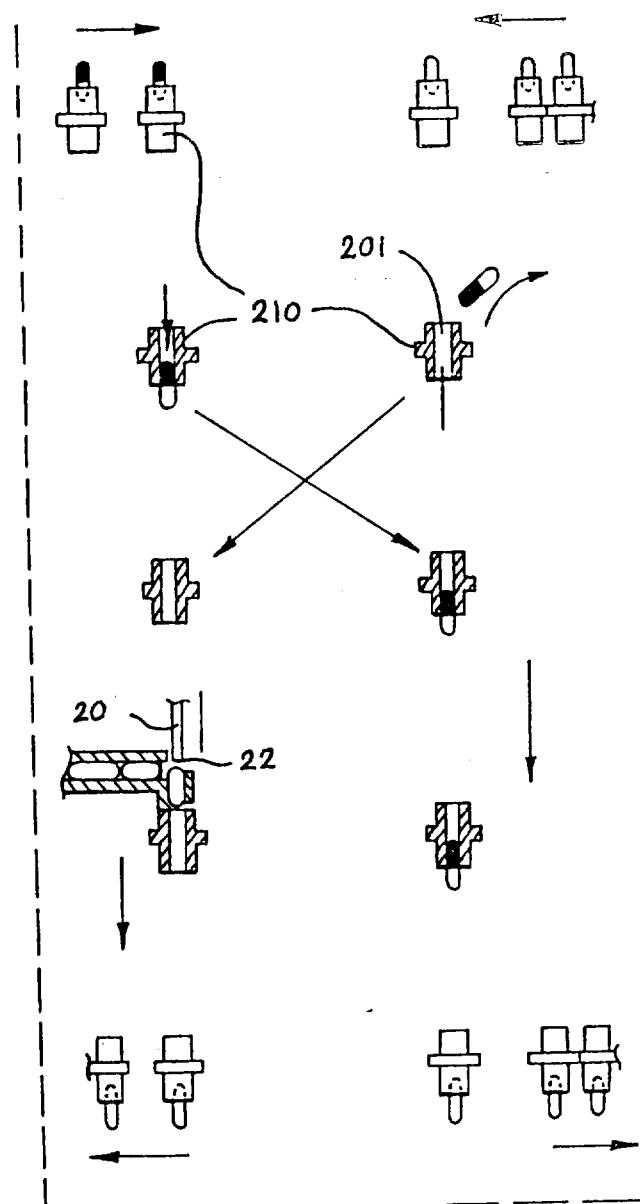


FIG. 1B

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FIG. 2

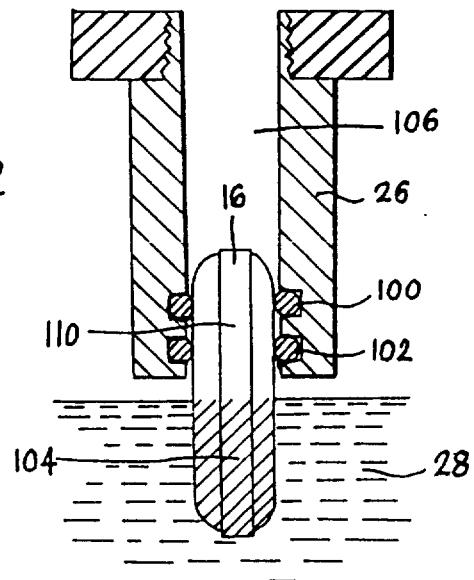
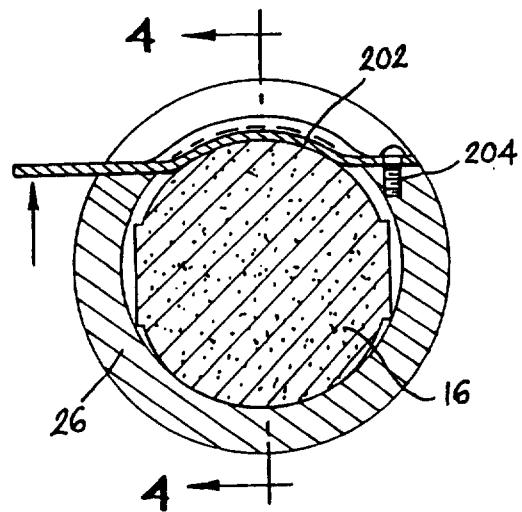


FIG. 3



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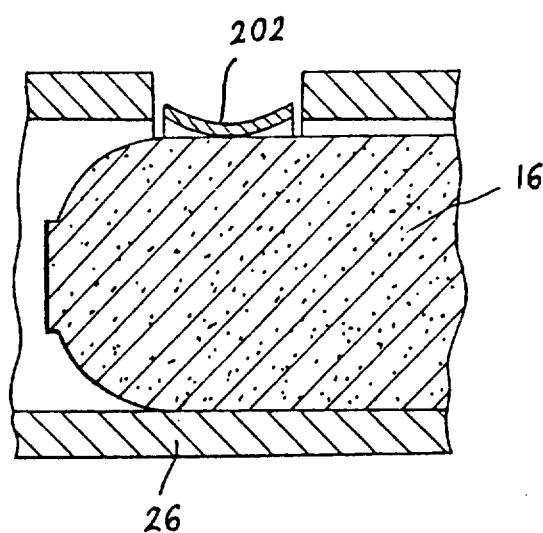


FIG. 4

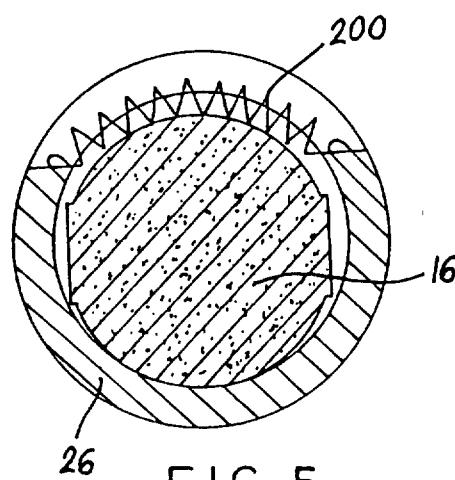


FIG. 5

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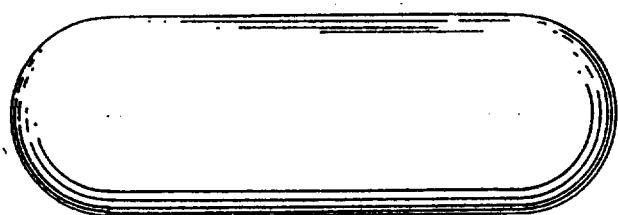


FIG. 6

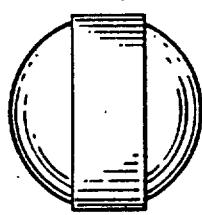


FIG. 7

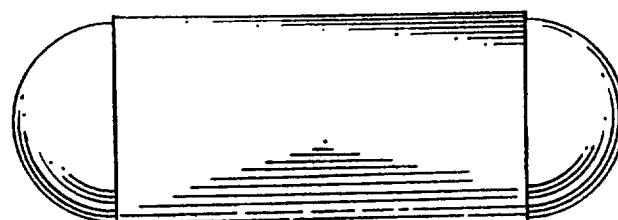


FIG. 9

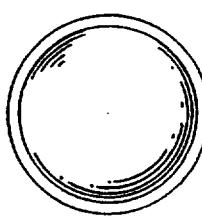


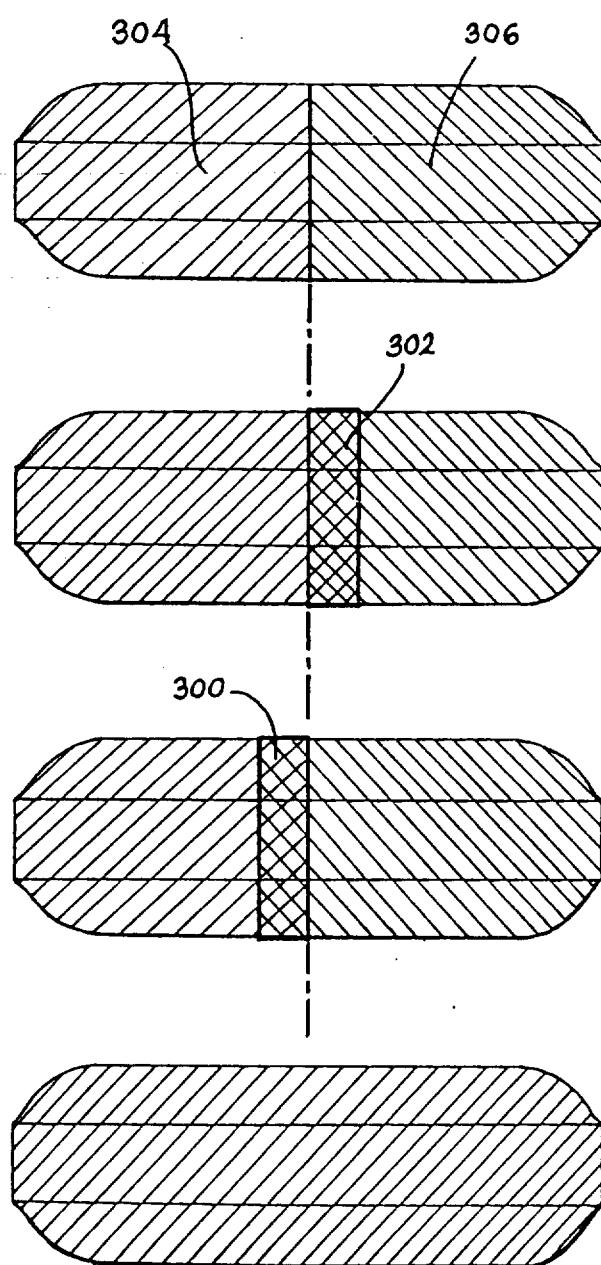
FIG. 10

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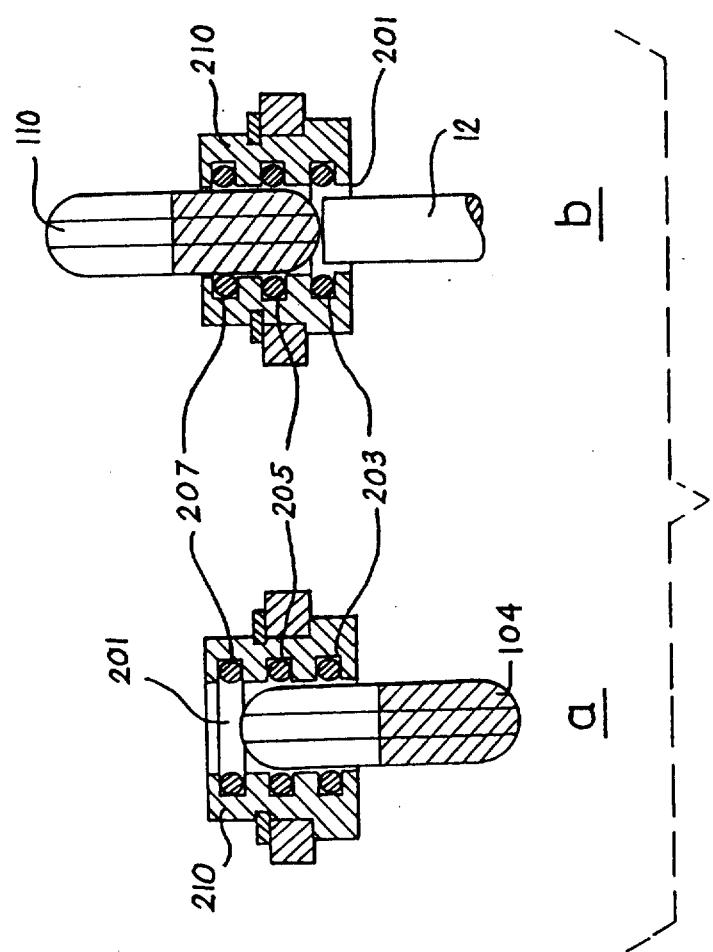
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>FIG. 8

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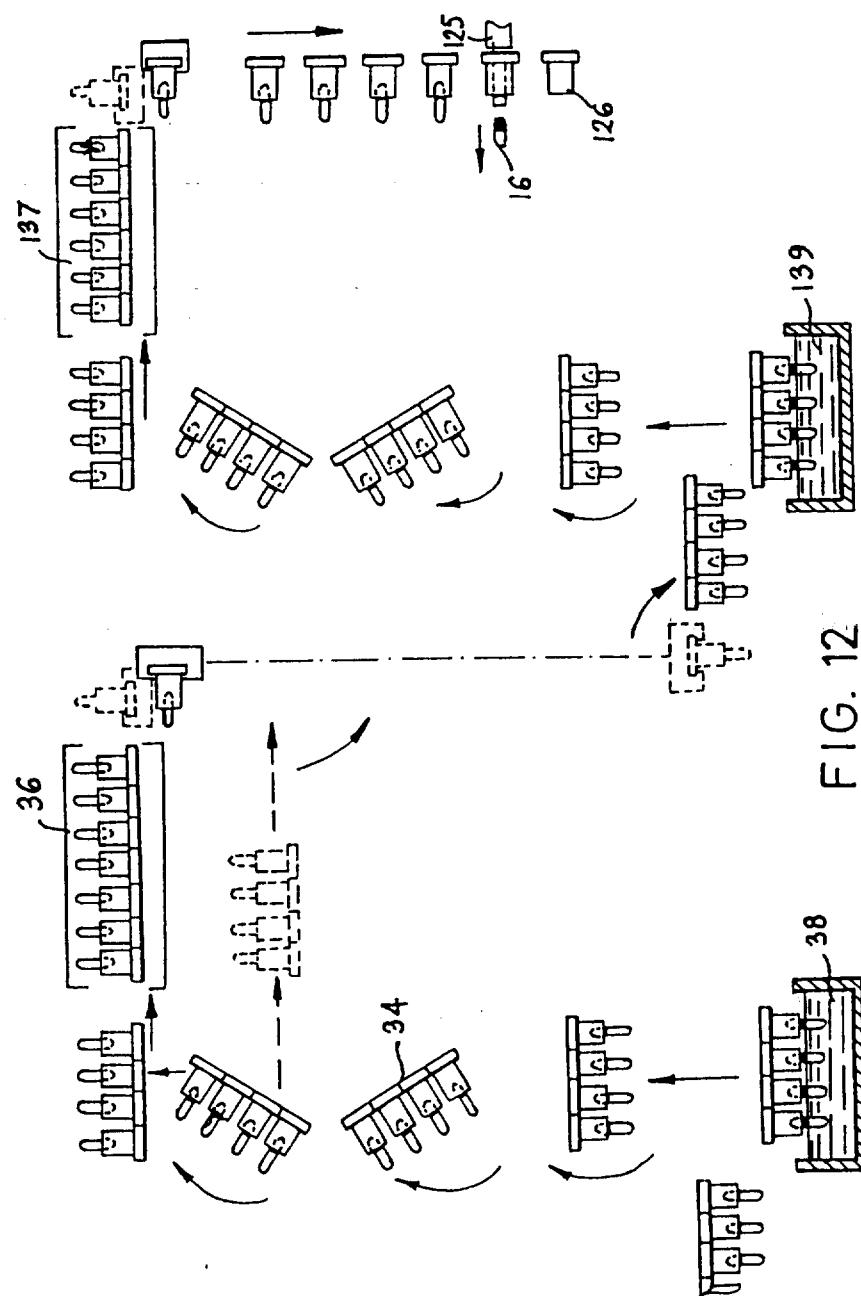


FIG. 12

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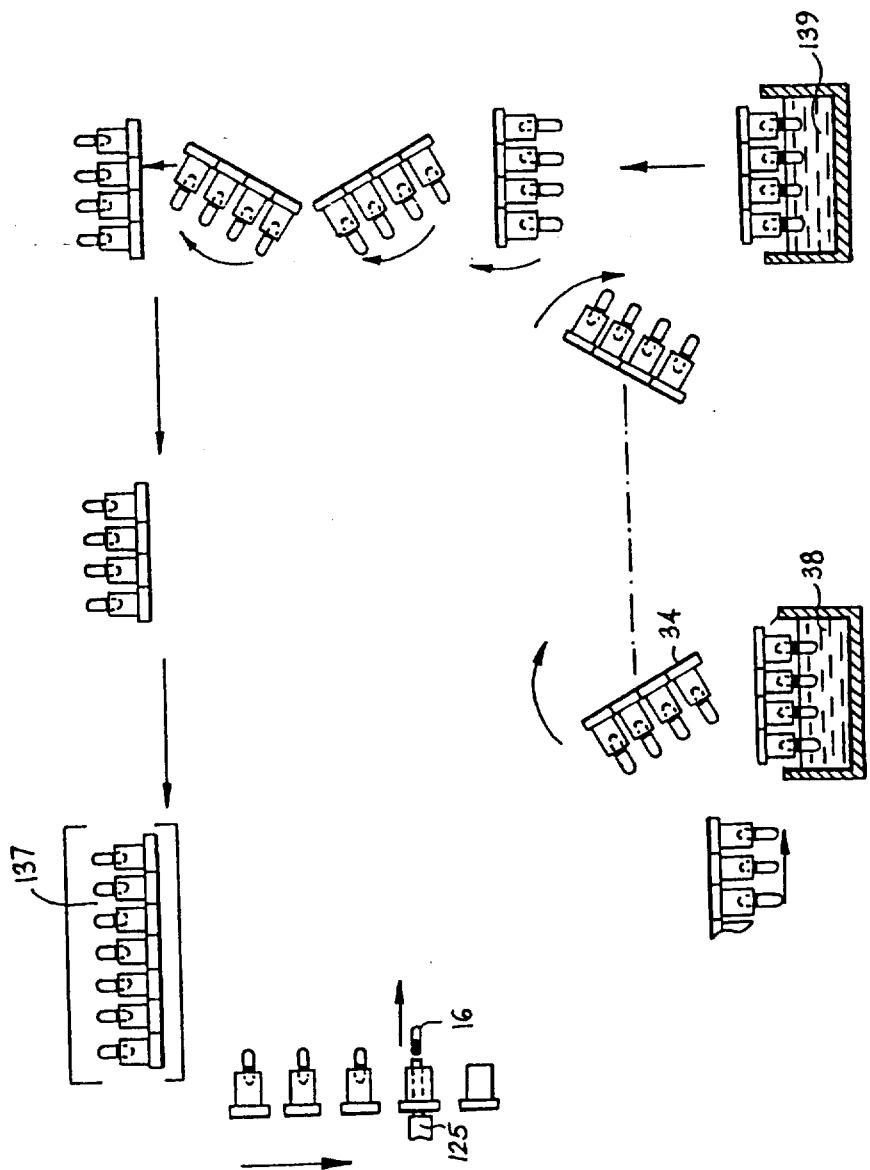


FIG. 13

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