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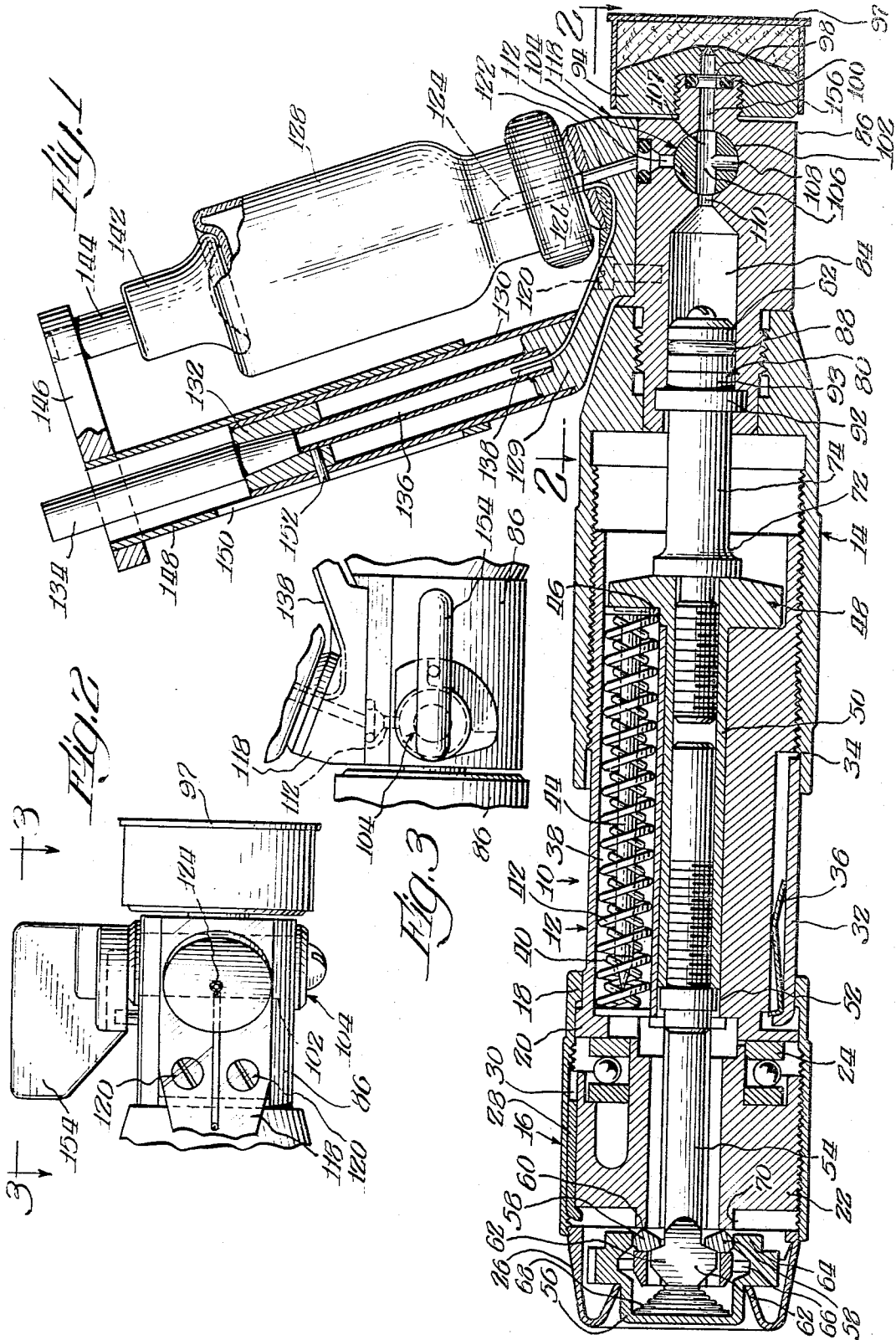
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HYPODERMIC JET INJECTOR

Filed Oct. 7, 1963

2 Sheets-Sheet 1



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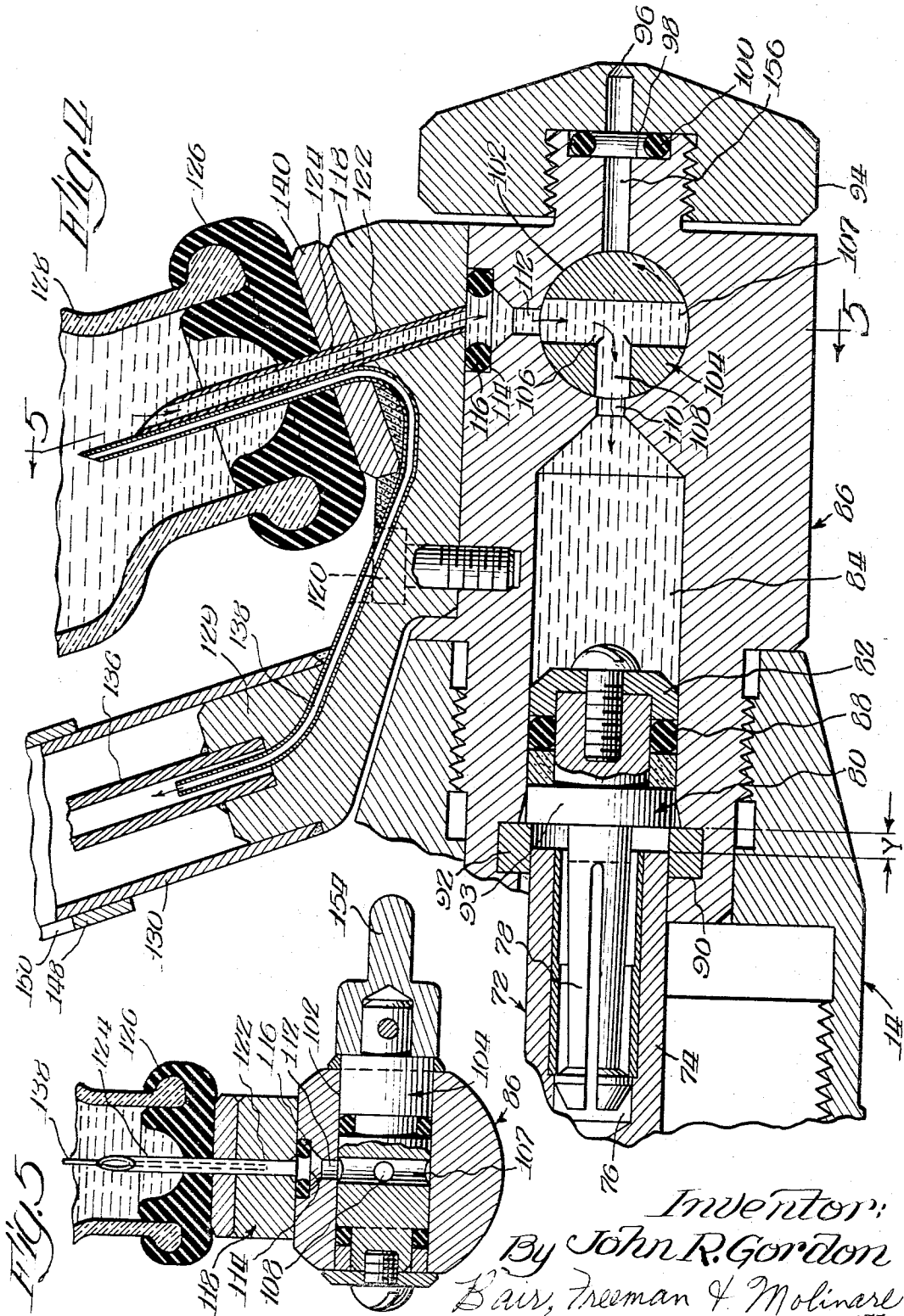
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Inventor:
By John R. Gordon
Bair, Freeman & Molinaro
Attys.

1

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HYPODERMIC JET INJECTOR

John R. Gordon, Grosse Pointe, Mich., assignor to

R. P. Scherer Corporation, Detroit, Mich.

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This invention relates to a manually operated hypodermic jet injector capable of administering successive injections of a liquid medicament through the epidermis. In some respects, this invention is similar to the multi-dose injector of Alfred W. Kath set forth in U.S. patent application Ser. No. 271,205, and in other respects it is similar to the single dose injector of Anthony Venditty, set forth in U.S. Patent No. 2,762,370.

Although the advantages of using hypodermic jet injectors rather than hypodermic needles are well-known, utilization by potential users, as by diabetics or pediatricians, is reduced due to relatively high costs of the medicament in ampule form or of the instrument itself. As a specific example, most diabetics continue to use hypodermic needles for administering insulin to themselves because of such high costs. In the case of a single-dose jet injector, similar to that set forth in U.S. Patent No. 2,762,370, ampules are used; although the instrument itself is relatively inexpensive, the required insulin dosage in ampule form costs approximately six times more than the cost of administering the insulin by a hypodermic needle. Also, in the case of a multi-dose jet injector, similar to that set forth in U.S. patent application Ser. No. 271,205, although the medicament is stored in a bottle in liquid form and not in ampule form, the cost of the instrument itself is quite high.

Furthermore, in the case of the multi-dose injectors, where the medicament is stored in the relatively inexpensive liquid form, the use of insulin presents a problem since in presently used long-acting insulin, such as Lente, PZI, and NPH, the insulin is mixed with a protein to provide a suspension, rather than a solution. In such suspensions, particles settle out of the suspension and are likely to cause clogging of the channels through which the insulin passes; also, there is a problem of the formation of crystals, which also cause clogging of the channels. Therefore, particularly in the case of insulin, it is highly important that the channels conducting insulin from the storage bottle or container to the medicament discharge chamber be constructed in such a manner that clogging thereof is substantially avoided.

It is therefore an important object of this invention to provide a hypodermic jet injector capable of storing a quantity of liquid medicament in bulk form while providing an injector of relatively inexpensive construction.

It is another object of this invention to provide a hypodermic jet injector which is mechanically loaded and operated for each injection and which is also capable of applying successive injections of medicament without separate loading of medicament for each injection.

It is also an object of this invention to provide a multi-dose jet injector particularly adapted for administering insulin.

It is still another object of this invention to provide a jet injector particularly adapted for the injection of insulin, wherein a quantity of insulin is stored in a bottle, and the channel from the bottle to the medicament discharge chamber is substantially straight, of minimum length and of substantially uniform cross-section, to thereby substantially reduce the possibility of clogging of the channel by particles settling out of the insulin suspension.

It is an additional object of this invention to provide a multi-dose jet injector particularly convenient to manipu-

2

late either by a person administering medicament to oneself or to another.

Further purposes and objects of this invention will appear as the specification proceeds.

5 A particular embodiment of the present invention is illustrated in the accompanying drawings wherein:

FIGURE 1 is a side elevational view of my hypodermic jet injector;

10 FIGURE 2 is a view taken along the line 2—2 of FIGURE 1;

FIGURE 3 is a view taken along the line 3—3 of FIGURE 2;

15 FIGURE 4 is an enlarged, partially sectioned view of the discharge end of the hypodermic jet injector of FIGURE 1; and

FIGURE 5 is a sectional view, on a reduced scale, taken along the line 5—5 of FIGURE 4.

Referring to the drawings, my hypodermic jet injector, generally indicated by the numeral 10, has the general appearance of an elongated body which includes a mechanism housing 12 that is enclosed at the forward end by a front sleeve 14, which is threadably connected thereto. At the opposite or rear end of the housing 12, a winding sleeve 16 is rotatably connected to the mechanism housing 12. The sleeve 16 includes an inturned flange 18, which co-operates with the annular ridge 20 of the housing 12 in order to prevent longitudinal movement between the housing 12 and the sleeve 16.

20 A latch housing 22, which also serves as a bearing support for thrust bearing 24, is threaded into the rear end of the winding sleeve 16. The thrust bearing 24 is interposed between the latch housing 22 and the mechanism housing 12 in order to permit relative rotation between the parts. A release button cap 26 is securely fastened to the winding sleeve 16 by means of a conventional non-reversible, bayonet type connection. The cap 26, the winding sleeve 16, and the latch housing 22 are all rotatable together, the sleeve 16 being locked to the housing 22 by means of a key 28 which fits into the slot 30 provided in the housing 22 and the sleeve 16.

35 The mechanism housing 12 is locked to the forward sleeve 14 in order to prevent rotation therebetween by means of a latch 32, which is adapted to fit into a series of longitudinal slots 34 provided in the inner surface of the sleeve 14. A spring 36 biases the latch 32 outwardly. The mechanism housing 12 includes a series of blind, axially extending openings 38 which are spaced circumferentially in the housing 12. Although single springs may be inserted into the openings 38, the springs are preferably used in pairs, the internal springs 40 being of a smaller diameter and of opposite helix to the external springs 42. Pins 44 are located concentrically of the springs 40 and 42 and extend through the openings 38 in order to prevent the buckling of the springs 40 and 42 within the openings 38. The forward ends of the springs seat in recesses 46 provided in the nut 48, the recesses 46 being concentric with the openings 38.

40 The nut 48 includes a sleeve-like extension 50, which projects rearwardly in the bore 52 of the housing 12; the extension 50 is prevented from rotational movement within the mechanism housing 12 by suitable means, as a key (not shown). The extension 50 is internally threaded and is adapted to threadably receive the lifting screw 54. Thus, upon rotation of the screw 54, the nut 48 is pulled rearwardly to thereby compress the springs 40 and 42 and load the injector 10.

45 A latch mechanism is provided for locking the lifting screw 54 against axial movement until such time as the springs 40 and 42 are released for injecting the medicament. Latch wedges 58 project through the openings 60 provided in the latch housing 22 and are adapted to en-

gage the fins 62 at the rear end of the lifting screw 54. The release button 56 includes an inwardly turned flange 64 which is adapted to bear against the outer ends of the latch wedges 58, to normally prevent them from retracting into the groove 66 of the latch housing 22. The release button 56, upon depression, causes the side walls to move forwardly into the groove 70 of the latch housing 22 against the compression of the spring 68, whereupon the groove 66 in the side wall of the button 56 becomes aligned with the openings 60 in the latch housing 22. Due to the slope on the bottom of the wedges 58, they move outwardly into the groove 66, to thereby release the lifting screw 54.

The internal threads at the upper end of the nut 48 threadably receive a ram 72 having an extended forward portion 74; the forward portion 74 is provided with a bore 76 which is adapted to slidably receive the split shank 78 of the plunger 80. The plunger head 82 is slidable within the medicament chamber 84 of the valve housing member 86. In order to prevent a back-flow of medicament, an O-ring 88 is provided on the plunger head 82 for sealing against the inner surfaces of the chamber 84. Also, a groove 90 is provided in the valve housing 86 in order to receive a stop ring 92 which is adapted to abut the flange 93 on the plunger 80 for preventing further rearward travel of the plunger 80 while permitting further rearward movement of the ram 72.

The valve housing member 86 is an elongated body and is adapted to threadably engage the forward end of the forward sleeve 14 and, at its opposite end, receives the discharge nose 94, which includes a medicament discharge orifice 96. The discharge nose is protectively covered by a cap 97. The forward end of the member 86 includes a recess 98 which is adapted to receive a gasket or O-ring 100 for providing a pressure tight seal between the discharge nose 94 and the member 86.

As illustrated most clearly in FIGURES 4 and 5 the member 86 includes a transverse aperture 102 which is adapted to rotatably receive a cylindrical valve member 104. Centrally of the valve member 104, there is provided a T-shaped opening 106 having a main channel 107 and a branch channel 108. The channel 108 is alignable with the bore 110 which is forward of the medicament chamber 84. The main channel 107, in one position, is adapted to be aligned with the inlet 112 in the upper end of the member 86, and in a 90° rotated position is alignable with both the bore 110 and the front port 156 in the housing 86 and is adapted to permit the flow of medicament from the medicament chamber 84 to the discharge orifice 96. A counter-bore 114 is provided above the inlet 112 for receiving an O-ring 116 in order to provide a seal between the member 86 and the bottle support member 118. The inlet 112 and the connecting counter-bore 114 form a cavity which interconnects the main channel 107 and channel defined in the needle 124 in a minimum straight line distance to further facilitate flow and prevent clogging and crystallization of the medicament.

As shown most clearly in FIGURE 5, the top side of the valve housing 86 includes a flattened upper surface for carrying the support member 118. The support 118 is secured to the flattened surface by means of screws 120 or other suitable fasteners. The bottle support member 118 includes an opening 122 which is adapted to support the straight hollow needle 124. The needle 124 is inserted into the cap 126 of the medicament bottle 128. In this regard, it is important to note that the hollow needle 124 provides a straight channel having a substantially uniform cross-section for the flow of medicament from the bottle 128 directly to the valve member 104, which communicates with the medicament chamber 84. Thus, the possibility of clogging or blocking the channel with crystals or particles, which drop out of suspension, is substantially eliminated since bends, turns, and restrictions, wherein clogging is likely to occur, are avoided. Also, the channel from the bottle 128 to the valve 104 is of mini-

mum length and this also contributes to the substantial elimination of the clogging of flow channels.

Of further importance, the bottle 128 is tipped rearwardly from its vertical axis in order that it does not interfere with the injecting operation, while at the same time is sufficiently forward so that it does not interfere with the winding or loading operation of the injector.

The bottle support member includes a rearwardly extending air filter support portion 129. A tubular member 130 is secured to the filter support portion 129 and interiorly carries a hollow plug 132. The plug 132, at its upper end, carries an air filter member 134. The lower end of the hollow plug 132 supports an air tube 136 which is located concentrically of the tube 130 and is received within an opening provided in the rear portion 129. A vent tube 138 is directed into the air tube 136 from the medicament bottle 128; the vent tube is located adjacent to the hollow needle 124 and opens into the bottle 128. Thus, as medicament is released, it is replaced by filtered air.

A resilient support pad 140 is interposed between the bottle support member 118 and the cap 126 of the bottle 128; furthermore, at the upper end of the bottle 128, a resilient support member 142 is carried on a shaft 144 which is mounted on an arm 146. The arm 146 in turn is fixed to the tube 148 which is slidably positioned around the tube 130. The tube 148 includes an elongated longitudinal slot 150 which is adapted to engage a rivet 150, which is secured to the hollow plug member 132 to thereby avoid rotation of the arm 146, while permitting axial movement thereof. Thus, by axial adjustment of the tube 148, the support member 148 may be positioned firmly against the bottom of the bottle 128, whereby the bottle is securely maintained in position between the resilient supports 140 and 142.

In the operation of the injector 10, assuming that medicament has just been discharged, the valve member 104 is rotated from the position of FIGURE 1 to the position of FIGURE 4, so that there is communication from the bottle 128 to the medicament chamber 84. The valve member 104 is rotated by an exterior handle 154 which is secured to it. The winding sleeve 16 is turned in a counterclockwise direction in order to move the lifting screw 54 rearwardly. As the end of the screw 54 contacts and compresses the spring 68, the release button 56 moves rearwardly and the wedges 58 are forced out of the groove 66. The wedges 58 then move underneath the fins 62 of the lifting screw 54 in order to latch the screw 54. At this time, the spring 68 snaps the release button 56 to the loading position.

The winding sleeve 16 is then rotated in a clockwise direction in order to turn the screw 54 in the sleeve 50 and to thereby move the nut 48 rearwardly and compress the springs 40 and 42. As the nut 48 moves rearwardly, the ram 72 and plunger 80 also move rearwardly, the movement of the plunger 80 being stopped by the ring 92. As the plunger 80 moves rearwardly, medicament flows from the bottle 126, through the straight channel 124 and the valve 104, and finally into the chamber 84; the flow of medicament is unobstructed by deposits in the channel 124. When the desired amount of loading is attained, as in the position of FIGURE 1, the winding is stopped and the valve 104 is rotated to the position of FIGURE 1 whereby the main channel is aligned with the medicament chamber 84 and the front port 156 in the valve housing 86. The injector 10 is then ready to inject the medicament. Upon pressing the button 56, the lifting screw 54 is moved forwardly, thereby moving the wedges 58 into the groove 66; this releases the lifting screw 54 as well as the nut 48, the ram 72 and the plunger 80. Also, the rearwardly tipped position of the bottle 126 contributes to ease of loading and also to ease of injecting since it is out of the way during both operations.

As shown most clearly in FIGURE 4, upon release of the lifting screw 54, the ram 72 moves forwardly a

5

short distance Y before there is actual movement of the plunger 80. The free movement of the ram 72 in this distance provides a high impact force in order to provide an initial high pressure sufficient to penetrate the epidermis and provide an opening through which the medicament may pass. Upon moving the distance Y, the impact force is dissipated, reducing the pressure at which the medicament is discharged through the epidermis.

It is clear from the foregoing that all of the objects have been accomplished. The injector 10 is of relatively inexpensive and simple construction and is particularly adaptable for use by diabetics who require periodic and frequent injections of insulin. The injector 10 combines advantages of a single-dose injector with advantages of a multi-dose injector since the relatively inexpensive construction of a single-dose injector is adapted for injections directly from a medicament bottle rather than from ampules. Furthermore, a straight and relatively short flow channel of substantially uniform cross-section is provided for the flow of medicament from the bottle to the valve member, thereby substantially avoiding clogging of the flow channel. This is particularly important in the case of insulin, which tends to precipitate solids which cause blocking of small openings. Also, the position of the medicament bottle is such that interference with the loading and injecting operations is substantially avoided.

While in the foregoing there has been provided a detailed description of a particular embodiment of the present invention, it is to be understood that all equivalents obvious to those having skill in the art are to be included within the scope of the invention as claimed.

What I claim and desire to secure by Letters Patent is:

1. A hypodermic jet injector comprising an injector body with a medicament discharge orifice at one end, a chamber for medicament in the body spaced from said orifice, a medicament container having a pierceable cap secured to said body, a valve in said body between said

6

chamber and discharge orifice for selectively directing medicament from the container to the chamber or communicating the chamber with said orifice and power means communicating with the chamber for selectively propelling medicament from the chamber through said orifice; said injector body having a flattened surface thereon above the valve, elongated bore means in said body perpendicular to said flattened surface communicating with said valve, a counter-bore at the outer end of said bore means, compressible seal means in said counterbore, a container-support having flat face means clamped against said flattened surface and said seal means, a straight tubular member carried by said container-support with one end of the tubular member opening directly into said bore means and the other end of the tubular member extending outwardly of the container-support, and a resilient pad on said container-support surrounding said tubular member as it emerges from said container-support, the capped end of said container being pierced by said tubular member and resting against said resilient pad.

2. A device as in claim 1 wherein the container-support is shaped to incline the resilient pad to face in a direction away from the end of the injector body at which the discharge orifice is located.

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RICHARD A. GAUDET, *Primary Examiner.*

ROBERT E. MORGAN, *Examiner.*