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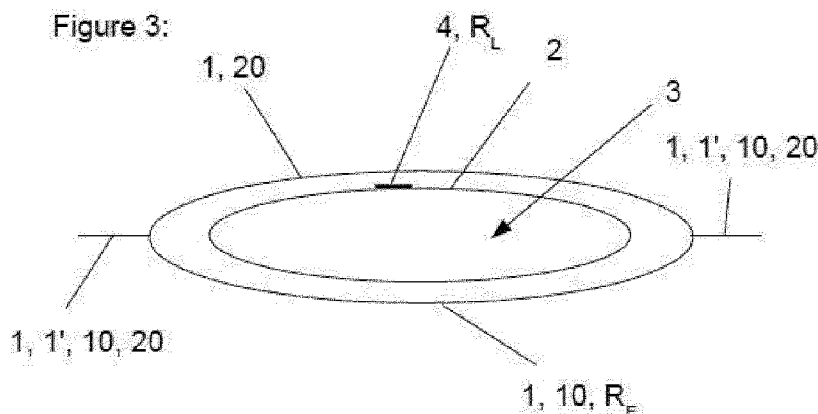
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(54) Title: CONTAINER CLOSURE SYSTEMS



(57) **Abstract:** The invention relates to a container closure system containing an overpouch with an intransparent first foil (10) and a transparent second foil (20), a transparent primary container (2) for holding a transparent pharmaceutical solution (3), wherein the transparent primary container (2) is packed within the overpouch (1) and labeled with at least one label (4) and wherein the at least one label (4) acts as a light absorbing segment having a reflection R_L for light in the range of 350 nm to 800 nm and an inner surface (18) of the intransparent first foil (10) of the overpouch (1) acts as a light reflecting background having a reflection R_F for light in the direction of the primary container (2) in the range of 350 nm to 800 nm with $R_F > R_L$. By the transparent second foil (20) and the inventive reflection properties it is achieved that the at least one label (4) on the primary container (2) is visible and readable. Additionally, visual inspection of the content of the transparent primary container (2) is possible. A good contrast is achieved to enhance machine and human readability.

Container closure system

DescriptionField of the invention:

The invention concerns a container closure system with a container which is filled with a pharmaceutical solution and with an overpouch in which the container is packed.

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Background of the invention:

Pharmaceutical solutions in containers and bags, respectively, often contain oxygen sensitive ingredients. As such a container it is usually used a primary bag which is fully transparent to allow visual inspection of the pharmaceutical solution in the primary bag. The primary bag usually could be made out of a multi-layer film and could have good chemical resistance, good welding characteristics and could be heat sterilisable. One or two tubes could be put in place by a heat welding process and serve to connect one or two ports to the multi-layer film. One port can be used for the infusion of the content of the primary bag into a patient. A second port can be used to inject additional compatible solutions into the content of the primary bag. Furthermore, labeling information is usually printed directly on the primary bag.

Such primary bags are often overwrapped by an overpouch, preferably immediately after filling and under reduced pressure or vacuum. Usually a secondary bag is used as such an overpouch. This outer secondary bag prevents largely gas permeation into the primary bag and serves therefore as a protection of the content of the primary bag against oxygen and water vapor transmission as well as against other environmental influences. Labels and/or barcodes are often provided on the secondary bag as well.

It is current praxis to use an aluminium overpouch as a secondary bag to provide a barrier for oxygen and water vapor. With a container closure system with such a secondary bag as an overpouch a visual inspection of the content of the primary bag is not possible. Furthermore, the label printed on the primary bag is covered by the intransparent overpouch. Thus, the label has to be reprinted on the overwrap because it is vital to have all necessary product information readily available without opening the overpouch.

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US 2013/0327677 A1 shows a transparent overwrap packaging system comprising a primary product contacting package, a secondary package comprising a multilayer high barrier oxygen overwrap. The primary product containing package can house detailed label instructions.

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It is therefore an object of the invention to provide an improved container closure system, in particular which makes a reprinting of the label of the primary bag onto the overpouch redundant. Another object of the invention is to allow or to improve visual inspection of the label and of the content of a transparent primary container of such a container closure system.

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Summary of the invention:

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The object of the invention is attained by a container closure system with the features of claim 1. Advantageous embodiments of the inventions are subject of the dependent claims.

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The inventive container closure system comprises the following components: An overpouch with an intransparent first foil and a transparent second foil and a transparent primary container, preferably embodied as a bag, for holding or storing a preferably transparent pharmaceutical solution. The transparent primary container is packed or encased within the overpouch and labeled with at least one label. According to a first alternative of the invention the at least one label acts as a light absorbing segment having a reflection R_L for light in the range of 350 nm to 800 nm and an inner surface of the intransparent first foil of the overpouch acts as a light reflecting background having a reflection R_F for light in the direction of the primary container in the range of 350 nm to 800 nm with $R_F > R_L$. According to a second alternative of the invention the at least one label acts as a light reflecting segment having a reflection R_L for light in the range of 350 nm to 800 nm and an inner surface of the intransparent first foil of the overpouch acts as a light absorbing background having a reflection R_F for light in the direction of the primary container in the range of 350 nm to 800 nm with $R_L > R_F$.

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By means of the transparent second foil and the inventive reflection properties it is achieved that the at least one label on the primary container is still visible and readable. Additionally, visual inspection of the content of the transparent primary container is possible. In particular by the the inventive reflection properties a good contrast is achieved to enhance machine and human readability of the label.

With this inventive container closure system design it is possible to overwrap a primary container which holds a pharmaceutical solution without covering labels which are arranged, especially printed, on the primary container. The primary container is filled with the pharmaceutical solution. The inventive container closure system is especially suitable for transparent pharmaceutical solutions. Therefore, the primary container is preferably filled with a transparent pharmaceutical solution. It is mentioned that under the term pharmaceutical solution not only liquid pharmaceutical solutions are meant but also solutions for infusion, nutrition and/or dialysis. This enumeration is exemplary only and not restricted to mentioned examples.

Preferably the entire area of the second foil is transparent and/or the entire area of the primary container is transparent. In case the area of the second foil is only partially transparent, the transparent area preferably comprises an area which is located above or covers the area of the primary container in which the label is located. Preferably the entire area of the first foil is intransparent. In case the area of the first foil is only partially intransparent, the intransparent area preferably comprises at least an area which is located below or covered by the area of the primary container in which the label is located.

Subsequently light is described as electromagnetic radiation as well. By the transparent second preferably multi-layer foil and the inventive reflection properties for electromagnetic radiation in the visible range of electromagnetic radiation between 350 nm to 800 nm of the foils of the overpouch and the primary container it is achieved that the labels on the primary container are still visible and are readable by machines.

In a first embodiment of the first alternative of the invention the label is provided by a dark color, preferably black color, and the inner surface of the intransparent first foil of the overpouch is provided by a light color, preferably white color. In a first embodiment of the second alternative of the invention the label is provided by a light color, preferably white color, and the inner surface of the intransparent first foil of the overpouch is provided by a dark color, preferably black color. An enhanced contrast and therefore an enhanced readability are achieved.

In a further embodiment of the first and the second alternative of the invention the label is imprinted on the outer side of the transparent primary container, preferably on the outer side of the transparent primary container facing the transparent second foil of the overpouch. Labels and/or barcodes could be printed on the primary bag using the hot stamp printing

technique. In this technique the ink is transferred from a carrier foil and melted to the surface of the bag during a short heating. This technique results in a print that is glossy and rub resistant and is even autoclavable.

- 5 In a further embodiment of the first and the second alternative of the invention the inner surface of the intransparent first foil of the overpouch is provided by a colored polymeric layer. Preferably the polymeric layer is made of or comprises polypropylene. In a preferred embodiment the color is part of the polymeric layer. The color belongs to the bulk. I.e. the color is a component of the blend to produce the polymer. The corresponding color can be
10 provided by pigments and/or by dyes. Preferably the color is not provided by an additional colored coating or painting on the polymeric layer surface.

The container closure system is characterized in a further embodiment such that the at least one label contains text information, a barcode, a data matrix, a symbol and/or a drawing.

- 15 Preferably it is or they are related to the content and/or the use of the primary container. The enumeration is exemplary and not restricted to the mentioned examples. The barcode can be a 1-dimensional or 2-dimensional barcode.

- The inventive container closure system is characterised in one embodiment of the first or the
20 second inventive alternative by the following reflection parameters: a) $0,5 \times R_F \geq R_L$ and $R_F \geq 0,5$ or b) $0,5 \times R_L \geq R_F$ and $R_L \geq 0,5$. Contrast and therefore human or machine readability are enhanced.

- In the case that the reflection R_F of the intransparent foil of the overpouch in the visible range
25 of electromagnetic radiation between 350 nm to 800 nm is at least 0,5 and at least twice as high as the reflection R_L of the at least one label on the primary container a good contrast of the at least one label of the primary container on the background of the intransparent foil of the overpouch is given. In the case that the reflection R_L of the at least one label on the primary container in the visible range of electromagnetic radiation between 350 nm to 800
30 nm is at least 0,5 and at least twice as high as the reflection R_F of the intransparent foil of the overpouch a good contrast of the at least one label of the primary container on the background of the intransparent foil of the overpouch is given as well.

- The difference of the reflected radiation in the visible range of electromagnetic radiation
35 between 350 nm to 800 nm gives a good contrast of the at least one label of the primary container on the background of the intransparent foil of the overpouch. Therefore it is ensured that human beings can read the at least one label as well as label reader machines

while the primary container is still packed in the overpouch. It is neither necessary to open the overpouch nor to reprint the at least one label on the overpouch to get the information of the label. The primary container with the pharmaceutical solution is safely packed into the overpouch and the information of the at least one label of the primary container is accessible at any time without opening the overpouch. Furthermore a visible inspection of the pharmaceutical solution within the primary container is possible by the inventive container closure system without opening the overpouch. The reflection of the transparent foils can be near zero and can therefore be neglected even if the radiation reflected by intransparent foil of the overpouch is running twice through it.

To get an even better contrast of the at least one label of the primary container on the background of the intransparent foil of the overpouch a symbol contrast is defined by the absolute value of the difference between the reflection R_F of the intransparent foil of the overpouch and the reflection R_L of the at least one label of the primary container in the visible range of electromagnetic radiation between 350 nm to 800 nm wherein this symbol contrast SC is specified by $SC = |R_F - R_L| \geq 0,5$. This feature of the reflection properties of the intransparent foil of the overpouch and the at least one label ensures a good machine readability without making too high and cost intensive demands on the optics of the machine which has to read the label. Preferably the parameters R_L , R_F and SC are determined according to test standard ISO/IEC15416.

Regarding a further embodiment of the inventive container closure system the reflection properties R_F and R_L of the intransparent foil of the overpouch and the label of the primary container in the visible range of electromagnetic radiation between 350 nm to 800 nm are specified as follows:

c) $R_F \geq 0,75$, preferred $R_F \geq 0,85$, especially preferred $R_F \geq 0,9$, and $R_L \leq 0,25$, preferred $R_L \leq 0,15$, especially preferred $R_L \leq 0,1$ or

d) $R_L \geq 0,75$, preferred $R_L \geq 0,85$, especially preferred $R_L \geq 0,9$, and $R_F \leq 0,25$, preferred $R_F \leq 0,15$, especially preferred $R_F \leq 0,1$.

These features ensure even a better machine readability since the contrast of the at least one label of the primary container on the background of the intransparent foil of the overpouch is furthermore increased.

The overpouch is a container or overpack for holding the primary container. The overpouch can be a blister-type container. In one embodiment of the invention the container closure system can be realized by an overpouch which has an intransparent first foil and a

transparent second foil which are weldable or welded together for carrying the primary container comprising a pharmaceutical solution.

Preferably the intransparent first foil and the transparent second foil are provided by a multilayer film. Preferably the intransparent first foil of the overpouch has an outer layer of a polyester layer or of a polypropylene layer and/or an inner layer of an intransparent polypropylene layer to provide the inner surface as the background. In one embodiment a metallic layer, preferably an aluminum layer is located between the inner layer and the outer layer.

Preferably the transparent second foil of the overpouch has an outer layer of polyester, preferably of polyethylene terephthalate, and an inner layer of polypropylene. In one embodiment an inorganic oxide layer is located between the inner layer and the outer layer.

The inorganic oxide layer of the transparent second multi-layer foil avoids the permeability of oxygen and water vapor. In particular by this inorganic oxide layer the object of the invention is attained that the labels of the primary container are readable while it is overwrapped and sealed by the sealed overpouch and while the impermeability of the overpouch for oxygen is still warranted. It is neither necessary to get the primary container out of the overpouch nor to reprint the at least one label on the overpouch to get the information of the label of the primary container. Furthermore, the inorganic oxide layer of the transparent second foil inhibits water vapor transmission and protects the primary container from any other environmental impact.

In a further embodiment of the invention the inorganic oxide layer of the transparent second foil is made of an oxide of aluminum and/or silicium, especially of an aluminum oxide of the form AlO_x . This oxide could be deposited directly on the surface of the polyethylene terephthalate layer of the second multi-layer foil so that no additional glue is necessary to get the oxide connected to the polyethylene terephthalate layer of the second foil of the overpouch. In case of aluminium foils for such an overpouch it is a multi-layer film preferably with a composition of more than 60 % polypropylene, more than 10 % aluminium, less than 20 % polyester and less than 5 % of a glue system (percentage by weight).

To simplify the manufacturing process of overwrapping and sealing the primary container within the overpouch the first and/or the second multi-layer foil are/is deepdrawable. By this embodiment it is possible that the form of the overpouch is adapted to the form of the primary container during the manufacturing process of a container closure system consisting of the

overpouch, the primary container and the pharmaceutical solution within the primary container.

The exemplary features of the layers of the intransparent first foil are as following:

- 5 - The polyester layer of the intransparent first foil of the overpouch consists of or comprises polyethylene terephthalate and/or the polypropylene layer of the intransparent first foil of the overpouch consists of or comprises oriented polypropylene.
- This polyester layer and/or this polypropylene layer and/or the metallic layer of the intransparent foil have a thickness between 5 μm and 50 μm , preferably 12 μm and 25 μm .
- 10 - The intransparent polypropylene layer of the intransparent first foil has a thickness between 50 μm and 150 μm , preferably 75 μm and 85 μm .

While the metallic layer, preferably aluminum layer, is responsible for the protection of oxygen, water vapor and light permeability of the intransparent foil of the overpouch, the intransparent polypropylene layer with the preferred thickness between 75 μm and 85 μm is responsible for a good and sufficient stiffness and mechanical stability and concomitantly being also a good water vapor and oxygen barrier of the overpouch.

To easily recognize labels and/or barcodes printed on the primary container, the intransparent polypropylene layer of the intransparent first foil is colored white in one embodiment. On such a white background labels as barcodes printed on the primary container are visible very well since they are usually printed with black or dark colored ink which provides a very good contrast to the white background. But it is still possible within the invention that the intransparent polypropylene layer of the intransparent first foil is colored dark, preferably black, while the labels are printed in bright color, preferably in white, on the primary container. In both cases a good contrast of the label of the primary container on the background of the intransparent foil of the overpouch is given.

The exemplary features of the layers of the transparent second foil are as following:

- 30 - In one embodiment the polyester layer, preferably a polyethylene terephthalate layer, of the transparent second foil of the overpouch has a thickness between 5 μm and 50 μm , preferably 12 μm and 25 μm .
- In one embodiment the polypropylene layer of the transparent second foil of the overpouch has a thickness between 50 μm and 150 μm , preferably 75 μm and 85 μm .

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Especially the polypropylene layer with the mentioned thickness raises the stiffness and mechanical stability of the overpouch once more. If the stiffness and/or mechanical stability

has to be particular high between the polyethylene terephthalate layer and the polypropylene layer of the transparent first foil of the overpouch, an additional polyester layer, preferably polyethylene terephthalate layer, can be located within these two layers. This layer can have a thickness between 5 μm and 50 μm , preferably 12 μm and 25 μm .

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On the outer wall of the transparent first foil of the overpouch the polyester layer, preferably the polyethylene terephthalate layer, is sealed with a heat sealable coating in a further embodiment. Overheating of the pharmaceutical liquid within the primary container during storage can be prevented.

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Since the intransparent first foil and/or the transparent second foil of the overpouch is/are preferably provided by single layers, theses single layers are laminated together by means of glue.

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The primary container of the inventive container closure system is preferable a fully transparent polyolefin bag, holding the (transparent) solution. The entire container closure system can be subjected to heat sterilization. The polypropylene layer of the intransparent first foil of the overpouch is white coloured. This assembly and especially the white background make the label printed directly with dark, respectively black, colour onto the primary container readable through the transparent second multi-layer foil of the overpouch. Hence, the second label printed usually on the overpouch becomes redundant. Furthermore, visual inspection can be performed more accurate on the white background. Therefore, only one label can be used for a double packed pharmaceutical liquid but still all information inevitably printed on the primary bag is disclosed and readable with the naked eye or with a machine, without a second label printed or glued on the overpouch. The human and/or machine readability of the label on the white background is significantly better as on the usually silver or dark background, for instance provided by an aluminium surface. In addition any potential change in quality can be detected without destruction of the overpouch.

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Especially for transparent primary containers which are filled with transparent pharmaceutical solutions like a paracetamol solution the containers can be furthermore inspected better with respect to quality parameters like colour change or visible particles. A colour change, often a sign of degradation of the finished product, is well detectable on the white background with the naked eyes or a machine. The same can be stated for visible particles. Both parameters can be tested without destruction of the overpouch, hence removal of the oxygen protecting shell.

The preferred method of sterilization of the container closure system is heat sterilization. In addition to the oxygen impermeable overpouch, an oxygen absorber is added between the primary container and the overpouch as a protecting agent against oxidation of the active pharmaceutical ingredient. The oxygen absorber, for example, could be positioned between two ports of the primary container, so that the readability of the label is not jeopardized. One port could be used for the infusion of the content of the primary bag into the patient. Another port, for example, could be used for injection (addition) of other compatible drugs. The primary container film could be a flexible multi-layer film made of polyolefine and has good chemical resistance, good welding characteristics, good water vapor barrier and is heat sterilisable.

If the entered oxygen in the overpouch cannot be bound anymore by the oxygen absorber or no absorber is present in the sealed overpouch within the sealed overpouch, an oxygen indicator can be present, which is preferable located on the outside of the primary container outside the area of the label of the primary container. Such an oxygen indicator changes its color if free oxygen is present so that it is easily recognizable if oxygen has entered the sealed overpouch and could not be bound by an oxygen absorber. This is important since the primary container comprises pharmaceutical solutions, which potentially contain an oxygen sensitive active ingredient.

The first and second multilayer foils of the overpouch exhibit an oxygen permeability of less than $3 \text{ cm}^3/(\text{m}^2 \cdot \text{day} \cdot \text{bar})$, in accordance to ISO standard 15105-2, at 23°C and 50 % r.h., whereby average values are measured around $0,4\text{-}0,5 \text{ cm}^3/(\text{m}^2 \cdot \text{day} \cdot \text{bar})$. The water vapor permeation is specified with $< 1 \text{ g}/(\text{m}^2 \cdot \text{day})$ in accordance to ISO standards 15106-3 at 23°C and 85 % r.h., whereby values of approximately $0,4 \text{ g}/(\text{m}^2 \cdot \text{day})$ are obtained. The foils were tested at conditions of $23^\circ\text{C}/85\%$ r.h after subjecting the foils to an autoclave cycle of $121^\circ\text{C}/30$ minutes.

The container closure system preferably comprises the primary container filled with the pharmaceutical solution representing a pharmaceutical product. The pharmaceutical solution can be a solution, a suspension or an emulsion comprising at least one compound and at least one liquid. The compound can be fully dissolved in the liquid, partially dissolved in the liquid, or suspended in the liquid. The compound of the pharmaceutical solution is for instance a therapeutic agent. Examples of therapeutic agents include, but are not limited to antiinfectives, anesthetics, analgesics, anticoagulants, chemotherapeutics, hormones, antihypertensives, antiinflammatories, antiemetics, bronchodilators, adrenergics, immunoglobulins, antipsychotics, antidepressants, and combinations thereof. In some

embodiments, the pharmaceutical solution comprises moxifloxacin, linezolid, levofloxacin, levetiracetam, vancomycin, cefepime, aztreonam, cefoxitin, ceftriaxone, cefazolin, cefotaxime, ceftazidime, gentamicin, oxacillin, nafcillin, penicillin, cefuroxime, ticarcillin, clavulanic acid, piperacillin, tazobactam, azithromycin, meropenem, ertapenem, tigecycline, micafungin, metronidazole, fluconazole, itraconazole, posaconazole, heparin, enoxaparin, dalteparin, theophylline, acetaminophen (paracetamol), ibuprofen, acetylcysteine, ropivacaine, lidocaine, propofol, decitabine, azacitidine, docetaxel, pemetrexed, palonosetron, aprepitant, fosaprepitant, famotidine, amiodarone, nitroglycerin, nicardipine, clevidipine, dobutamine, magnesium sulfate, sodium chloride, potassium chloride, lactated ringer's, dextrose, mannitol, or a combination thereof. The pharmaceutical solution can contain one or more excipients. Non-limiting examples of excipients include carriers, diluents, salts, buffers, stabilizers, solubilizers, preservatives, chelating agents, antioxidants, and tonicity contributors.

Further goals, advantages, features and applications of the invention arise from the following description of embodiments of the invention on the basis of the figures. The features of the different embodiments are able to be combined with one another. Thereby all features described and all features shown in the figures alone or in arbitrary reasonable combination provide the subject matter of the invention independent of their conclusion in the claims or their dependency.

Brief description of the drawings:

They show:

Figure 1: an exploded view of a schematically sectional view of one embodiment of a transparent foil of an overpouch of an inventive closure system,

Figure 2: an exploded view of a schematically sectional view of one embodiment of an intransparent foil of an overpouch inventive of an inventive closure system,

Figure 3: one embodiment of an inventive container closure system in a schematically sectional view,

Figure 4: the inventive container closure system of figure 3 in a view from above through the transparent foil of an overpouch of the inventive container closure system,

Figure 5: one further embodiment of an inventive container closure system in a schematically sectional view and

- 5 Figure 6: the inventive container closure system of figure 5 in a view from above through the transparent foil of an overpouch of the inventive container closure system.

Subsequently, preferred but exemplary embodiments of the invention are described in more
10 detail with regard to the figures.

Detailed description of the invention:

15 In the figures 1 and 2 a transparent foil 20 and an intransparent foil 10 of one embodiment of an overpouch 1 of an inventive container closure system are shown in an exploded view of a schematically sectional view. The sectional composition of the transparent foil 20 and the intransparent foil 10 of the overpouch 1 are clearly visible.

20 The transparent second foil 20 of the overpouch 1 which is transparent is shown in figure 1. In one embodiment the transparent foil 20 is not deepdrawable. To the outside of the overpouch 1 the transparent foil 20 is delimited by a polyethylene terephthalate layer 21 which is used as an outer wall 29 of the transparent foil 20. In the shown embodiment the polyethylene terephthalate layer 21 is coated on the outside with a heat sealable coating 24
25 which inhibits heat transmission into the overpouch 1 and into a primary container 2 of a container closure system, respectively, when the filled primary container 2 is sealed by the overpouch 1 and, for example, stored in a storage.

On the inner side of the polyethylene terephthalate layer 21 an inorganic oxide layer 23, in
30 particular an aluminum oxide layer, is preferably directly deposited. This oxide layer 23 builds a barrier for oxygen, water vapor and other gases within the transparent foil 20.

The polyethylene terephthalate layer 21, the heat sealable coating 24 and the inorganic oxide layer 25 form a layer assembly having a thickness of about 10 μm to 15 μm , preferably
35 12 μm . This layer assembly is bonded to an additional polyethylene terephthalate layer 25, preferably having a thickness between 12 μm and 25 μm , by means of glue 31. This additional polyethylene terephthalate layer 25 is furthermore bonded to a transparent

polypropylene layer 22, preferably having a thickness between 75 μm and 85 μm , by means of glue 30. The main function of the additional polyethylene terephthalate layer 25 and the polypropylene layer 22 is in particular to enhance the stiffness and the mechanical stability of the transparent foil 20, the overpouch 1 and the container closure system. Furthermore, the transparent polypropylene layer 22 forms the inner wall 28 of the second multi-layer foil 20 of the overpouch 1. Preferably all layers of the transparent foil 20 are transparent or essentially transparent.

If a primary container 2 which is filled with a pharmaceutical solution 3 is sealed with an overpouch 1 containing a transparent foil 20 as the above described second multi-layer foil 20, it is possible to look through the transparent foil 20 into the interior of the overpouch 1 and to recognize label 4, for instance in form of barcodes 4a arranged on the primary container, while gas permeation through transparent foil 20 is inhibited essentially by the inorganic oxide layer 23. So it is possible to read all information labeled on the primary container 1 without destroying the overpouch 1 and the protection for the primary container 1 and the pharmaceutical solution therein, respectively.

The first multi-layer foil 10 of the overpouch 1 which is intransparent is shown in figure 2. In a preferred embodiment the intransparent first multi-layer foil 10 is deepdrawable. To the outside of the overpouch 1, the intransparent foil 10 is delimited in this embodiment by a polyester layer 11 which is used as an outer wall 19 of the intransparent foil 10. In a specific embodiment, the polyester layer 11 is formed out of polyethylene terephthalate, preferably having a thickness between 12 μm and 25 μm . In a further embodiment the intransparent foil 10 is delimited by a polypropylene layer 11 which is used as an outer wall 19 of the intransparent foil 10. In a specific embodiment, the polypropylene layer 11 is formed out of oriented polypropylene, preferably having a thickness between 12 μm and 25 μm .

This layer 11 is bonded to an aluminum layer 13, preferably having a thickness between 12 μm and 25 μm , by means of glue 32. That aluminum layer 13 builds a barrier for oxygen, water vapor and other gases within the intransparent foil 10.

Furthermore, this aluminum layer 13 is bonded to a polypropylene layer 12, preferably having a thickness between 75 μm and 85 μm , by means of glue 33. One function of this polypropylene layer 12 is to enhance the stiffness and the mechanical stability of the intransparent foil 10, the overpouch 1 and the container closure system. The main function of this polypropylene layer 12 is hidden in its white color. Because of the white coloring of the polypropylene layer 12 labels 4 as barcodes 4a which are printed in black or generally in

dark color on transparent primary containers 2 which are filled with a preferable transparent pharmaceutical solutions 3 are very good readable by humans and machines. The black or dark printing on the primary container gives a good contrast to the white background.

Furthermore, because of the aluminum layer 13 of the intransparent foil 10 and the inorganic oxide layer 23 of the second multi-layer foil 20, a good protection against oxygen, water vapor and other gases is provided for pharmaceutical solutions 3 within a primary container 2 when the overpouch 1 with the described transparent foil 20 and the intransparent foil 10 are used for overwrapping and sealing.

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Figure 3 shows one embodiment of an inventive container closure system in a schematically sectional view wherein a primary container 2 with a pharmaceutical solution is overwrapped and sealed by an inventive overpouch 1. The filled primary container 2 is securely held within the overpouch 1 and therefore, the pharmaceutical solution 3 is protected against oxygen, water vapor and other gases. On the primary container 2 beneath the transparent foil 20 of the overpouch 1 a label 4 is printed. The label 4 can be embodied as and/or can comprise a 1-dimensional or 2-dimensional barcode or data matrix. As clearly shown in figure 3, the overpouch 1 is formed out of the intransparent foil 10 and the transparent foil 20 which are welded together in a welding area 1' in a boundary area of the overpouch 1.

20

If a machine or a human being wants to read the label 4 of the primary container 2 it is necessary that electromagnetic radiation has to fall onto the container closure system. Since human beings can see electromagnetic radiation in the visible range between 350 nm to 800 nm the label 4 of the primary container 2 has a reflection property of R_L while the intransparent foil 20 of the overpouch 1 has a reflection property of R_F in that range. For instance in this embodiment the following parameters could be used: $R_F \leq 0,05$ while $R_L \geq 0,80$. The difference of the reflected radiation in the visible range of electromagnetic radiation between 350 nm to 800 nm gives a good contrast of the at least one label of the primary container on the background of the intransparent foil of the overpouch since a symbol contrast SC which is defined by the absolute value of the difference of R_F and R_L has a high value of at least 0,75. Therefore the contrast of the label 4 of the primary container 2 is very good on the background of the intransparent foil 20 of the overpouch so that the label can be read easily by human beings as well as by machines through the transparent foil 10 of the overpouch.

35

The reflection of the transparent foil 20 of the overpouch 1 and the transparent primary container 2 and the transparent pharmaceutical solution 3 do not essentially contribute and

can therefore be neglected even if the radiation reflected by intransparent foil 10 of the overpouch 1 is running twice through it.

Figure 4 shows one embodiment of an inventive container closure system in a view from above through the transparent second multi-layer 20 foil of an overpouch 1 of the container closure system. In particular, this illustration makes the goal of the invention very clear.

The primary container 2 is filled with a pharmaceutical solution 3 and printed with a label 4, 4b, 4c which contains for instance information of the content 6 and information of the use 7 of the primary container. By reading this information 6 and 7 it is possible that humans can be informed directly about the use and the content. Furthermore, a label 4 with a barcode 4a was printed on the primary container 2 so that all necessary information is stored therein and can be read by a machine which can deliver this information to a data management system, especially a healthcare, patient and/or drug management and administration system. The labels 4 and the barcode 4a are printed preferably with black color, for instance ink, on the transparent primary container 2 so that this black ink builds up a very good contrast to the white intransparent polypropylene layer 12 of the first multi-layer foil 10 of the overpouch 1 which is visible because of the intransparent foil 20 of the overpouch and the different reflections R_F of the intransparent foil 10 of the overpouch 1 and R_L of the labels 4, 4a, 4b and 4c.

Especially when the pharmaceutical solution 3 within the primary container 2 is transparent, too, it is even possible to inspect the pharmaceutical solution 3 optically by humans or by machines. On the white background contaminations especially in form of particles or turbidities or color changing have a good visibility and are good indicators or signs of degeneration of the pharmaceutical solution 3 within the primary container 2.

Additionally, ports 26, 27, oxygen absorber 8 and oxygen indicator 9 are illustrated in figure 4. The oxygen absorber 8 is located on the surface of the primary container 2 in direction to the overpouch 1. This oxygen absorber 8 is located between two ports 26 and 27 of the primary container so that it does not block the labels 4 or the barcode 4a so that they are still visible. One port 26 can be used for infusion of a pharmaceutical solution 3 while the other port 27 can be used for adding additional pharmaceuticals or drugs into the pharmaceutical solution after destroying the overpouch 1 and before infusing the pharmaceutical solution into a patient.

Although the overpouch 1 should be gas tight, it is possible that leaks occur through which, in particular, oxygen can enter the overpouch and contaminate the pharmaceutical solution 3 within the primary container 2. To detect such an entry of oxygen into the sealed overpouch an oxygen indicator 9 is also located on the surface of the primary container 2 in a way that
5 neither the labels 4 nor the barcode 4a are blocked. Such an oxygen indicator 9 changes its color if oxygen is present in the sealed overpouch 1 so that an oxygen entry easily can be determined.

As already mentioned before, figures 3 and 4 illustrate an embodiment where the label 4 is
10 provided by black color on the transparent primary container 2. The inner surface 18 of the intransparent first foil 10 is provided by white color, preferably a white colored layer.

Finally, figures 5 and 6 illustrate a further embodiment of an inventive container closure system. In contrast to figure 3 and 4 the label 4 is provided by white color on the transparent
15 primary container 2. The inner surface 18 of the intransparent first foil 10 is provided by black color, preferably a black colored layer. In addition, it is illustrated that only a part of the inner surface 18 of the intransparent first foil 10 is provided with the black color. The intransparent area is located below or covered by the area of the primary container 2 in which the label 4 is located.

20 It will be understood that the invention may be embodied in other specific forms without departing from the spirit or central characteristics thereof. The present examples and embodiments, therefore, are to be considered in all respects as illustrative and not restrictive, and the invention is not to be limited to the details given herein. Accordingly, features of the
25 above described specific embodiments can be combined with one another. Further, features described in the summary of the invention can be combined with one another. Furthermore, features of the above described specific embodiments and features described in the summary of the invention can be combined with one another.

Reference signs

	1	overpouch
	1'	welded area of the overpouch
5	2	primary container
	3	pharmaceutical solution
	4	label
	4a	barcode
	4b	label with information of the content of the primary container
10	4c	label with information of the use of the primary container
	6	information of the content of the primary container
	7	information of the use of the primary container
	8	oxygen absorber
	9	oxygen indicator
15	10	intransparent foil
	11	polyester layer or polypropylene layer
	12	intransparent colored polypropylene layer
	13	aluminum layer
	18	inner wall or surface
20	19	outer wall or surface
	20	transparent foil
	21	polyethylene terephthalate layer
	22	polypropylene layer
	23	inorganic oxide layer
25	24	heat sealable coating
	25	polyethylene terephthalate layer
	26	port
	27	port
	28	inner wall
30	29	outer wall
	30	glue
	31	glue
	32	glue
	33	glue
35	R _F	reflection of the intransparent foil 10 or its inner surface 18
	R _L	reflection of the label 4, 4a, 4b and 4c

Claims:

1. Container closure system containing
an overpouch (1) with an intransparent first foil (10) and a transparent second foil
5 (20),
a transparent primary container (2), preferably a bag (2), for holding a transparent
pharmaceutical solution (3),
wherein the transparent primary container (2) is packed within the overpouch (1) and
labeled with at least one label (4) and
10 wherein the at least one label (4) acts as a light absorbing segment having a
reflection R_L for light in the range of 350 nm to 800 nm and an inner surface (18) of the
intransparent first foil (10) of the overpouch (1) acts as a light reflecting background having a
reflection R_F for light in the direction of the primary container (2) in the range of 350 nm to
800 nm with $R_F > R_L$ or
15 wherein the at least one label (4) acts as a light reflecting segment having a reflection
 R_L for light in the range of 350 nm to 800 nm and an inner surface (18) of the intransparent
first foil (10) of the overpouch (1) acts as a light absorbing background having a reflection R_F
for light in the direction of the primary container (2) in the range of 350 nm to 800 nm with R_L
> R_F .
20
2. Container closure system according to claim 1 characterized in
that the label (4) is provided by a dark color, preferably black color, and the inner
surface (18) of the intransparent first foil (10) of the overpouch (1) is provided by a light color,
preferably white color, or
25 that the label (4) is provided by a light color, preferably white color, and the inner
surface (18) of the intransparent first foil (10) of the overpouch (1) is provided by a dark color,
preferably black color.
3. Container closure system according to one of the preceding claims characterized in
30 that the label (4) is imprinted on the outer side of the transparent primary container
(2), preferably facing the transparent second foil (20) of the overpouch (1), and/or
that the inner surface (18) of the intransparent first foil (10) of the overpouch (1) is
provided by a colored polymeric layer, preferably made of or comprising polypropylene.
- 35 4. Container closure system according to one of the preceding claims characterized in
that the at least one label (4) contains text information (4b, 4c), a barcode (4a), a symbol
and/or a drawing, in particular related to the content and/or the use of the primary container
(2).

5. Container closure system according to one of the preceding claims characterized in that

a) $0,5 \times R_F \geq R_L$ and $R_F \geq 0,5$

5 or

b) $0,5 \times R_L \geq R_F$ and $R_L \geq 0,5$

and preferably characterized by a symbol contrast SC wherein $SC = |R_F - R_L| \geq 0,5$.

6. Container closure system according to claim 5 characterized in that

10 c) $R_F \geq 0,75$, preferred $R_F \geq 0,85$, especially preferred $R_F \geq 0,9$, and $R_L \leq 0,25$, preferred $R_L \leq 0,15$, especially preferred $R_L \leq 0,1$ or

d) $R_L \geq 0,75$, preferred $R_L \geq 0,85$, especially preferred $R_L \geq 0,9$, and $R_F \leq 0,25$, preferred $R_F \leq 0,15$, especially preferred $R_F \leq 0,1$.

15 7. Container closure system according to one of the preceding claims characterized in that the intransparent first foil (10) of the overpouch (1) is a multilayer-foil having an outer layer (11) and a, preferably intransparent, inner layer (12) providing the inner surface (18).

8. Container closure system according to claim 7 characterized in

20 that a metallic layer (13), preferably an aluminum layer (13), is located between the outer layer (11) and the inner layer (12) of the intransparent first foil (10) and/or

that the outer layer (11) of the intransparent foil (10) is a polyester layer (11) or a polypropylene layer (11) and/or that the inner layer (12) of the intransparent first foil (10) is an intransparent polypropylene layer (12).

25

9. Container closure system according to claim 8 characterized in that the polyester layer (11) of the intransparent first foil (10) is a polyethylene terephthalate layer (11) or the polypropylene layer (11) of the intransparent first foil (10) is an oriented polypropylene layer (11).

30

10. Container closure system according to one of the claims 8 to 9 characterized in that the polyester layer (11), the polypropylene layer (11) and/or the metallic layer (13) of the intransparent first foil (10) have a thickness between 5 μm and 50 μm and/or the intransparent polypropylene layer (12) of the intransparent first foil (10) has a thickness

35 between 50 μm and 150 μm and/or

that the layers (11, 13, 12) of the intransparent first foil (10) of the overpouch (1) are laminated together by a glue (32, 33).

11. Container closure system according to one of the preceding claims characterized in that the transparent second foil (20) of the overpouch (1) is a multilayer foil having an outer layer (21, 24) and an inner layer (22).

12. Container closure system according to claim 11 characterized in that an inorganic oxide layer (23), preferably made of an oxide of aluminum or silicium, is located between the outer layer (21, 24) and the inner layer (22) of the transparent second foil (20) and/or

that the outer layer (21) of the transparent second foil (20) is a polyester layer, preferably a polyethylene terephthalate layer (21), and/or the inner layer (22) is a polypropylene layer (22).

13. Container closure system according to one of the claims 11 to 12 characterized in that the polyester layer (21), preferably the polyethylene terephthalate layer (21), of the transparent second foil (20) is sealed on the outside of the overpouch (1) with a heat sealable coating (24) and/or

that the polyester layer (21), preferably the polyethylene terephthalate layer (21), of the transparent second foil (20) of the overpouch (1) has a thickness between 5 μm and 50 μm and/or the polypropylene layer (22) of the transparent foil (20) of the overpouch (1) has a thickness between 50 μm and 150 μm .

14. Container closure system according to one of the claims 11 to 13 characterized in that an additional polyester layer (25), preferably a polyethylene terephthalate layer (25), in particular having a thickness between 5 μm and 50 μm , is located between the polyester layer (21), preferably the polyethylene terephthalate layer (21) and the polypropylene layer (22) of the transparent second foil (20) of the overpouch (1) and/or

that the layers (21, 25, 22) of the transparent second foil (20) are laminated together by a glue (30, 31).

15. Container closure system according to one of the preceding claims characterized in that the sealed overpouch (1) contains an oxygen absorber (8) and/or an oxygen indicator (9), preferably located outside of the transparent primary container (2) and/or outside of the area of the at least one label (4) of the transparent primary container (2) and/or

that the intransparent first foil (10) is deepdrawable and/or the transparent second foil (20) of the overpouch (1) is not deepdrawable and/or that no additional label is placed, preferably imprinted, on the overpouch (1) and/or

that the primary container (2) is filled with the pharmaceutical solution (3).

Figure 1:

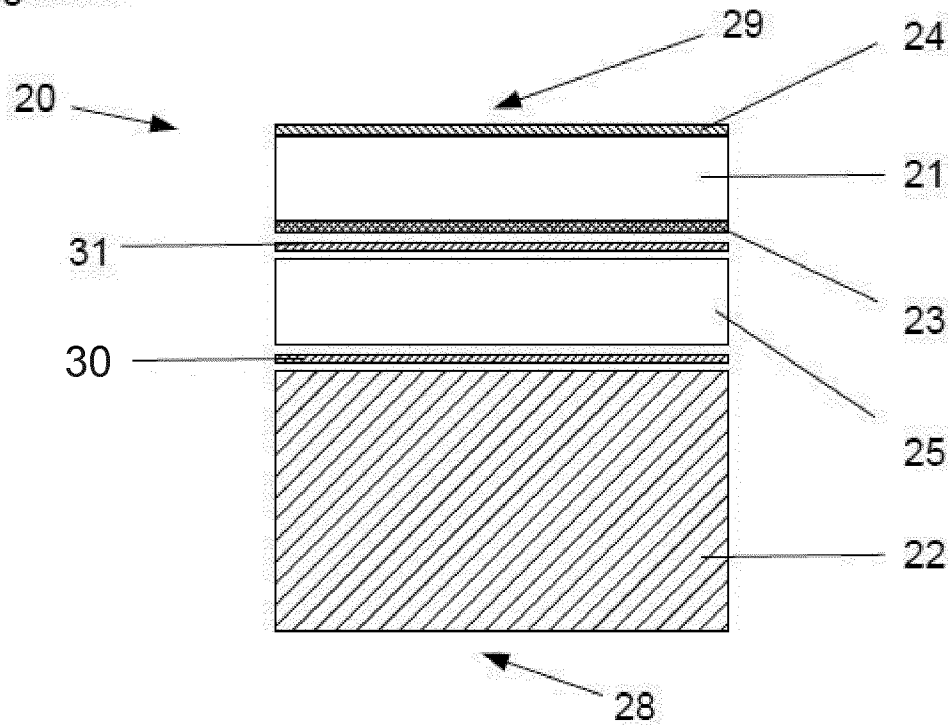


Figure 2:

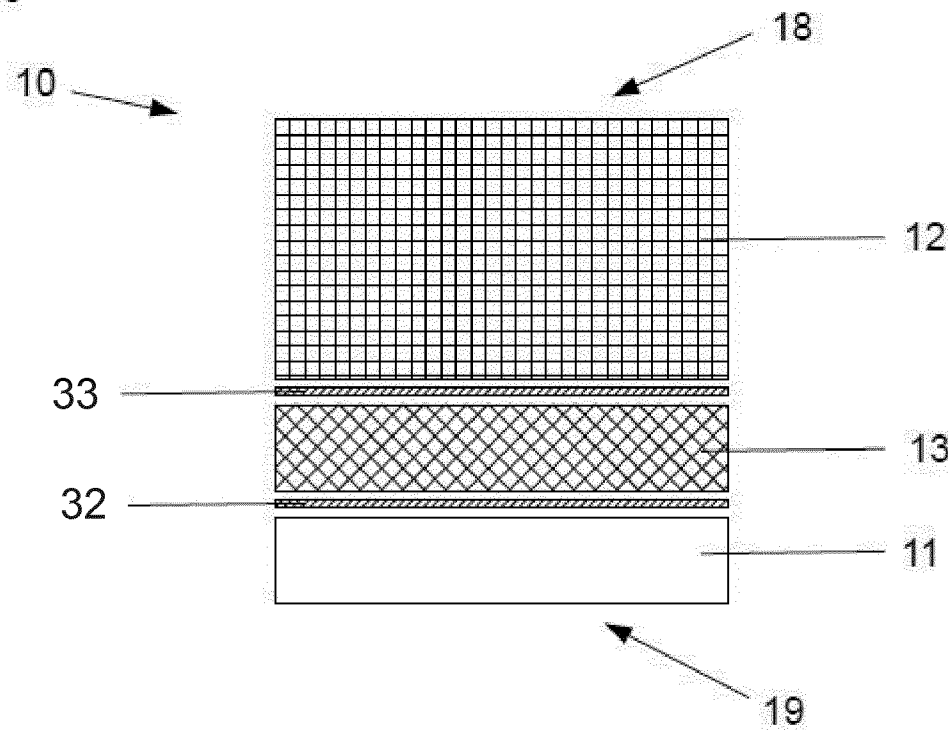


Figure 3:

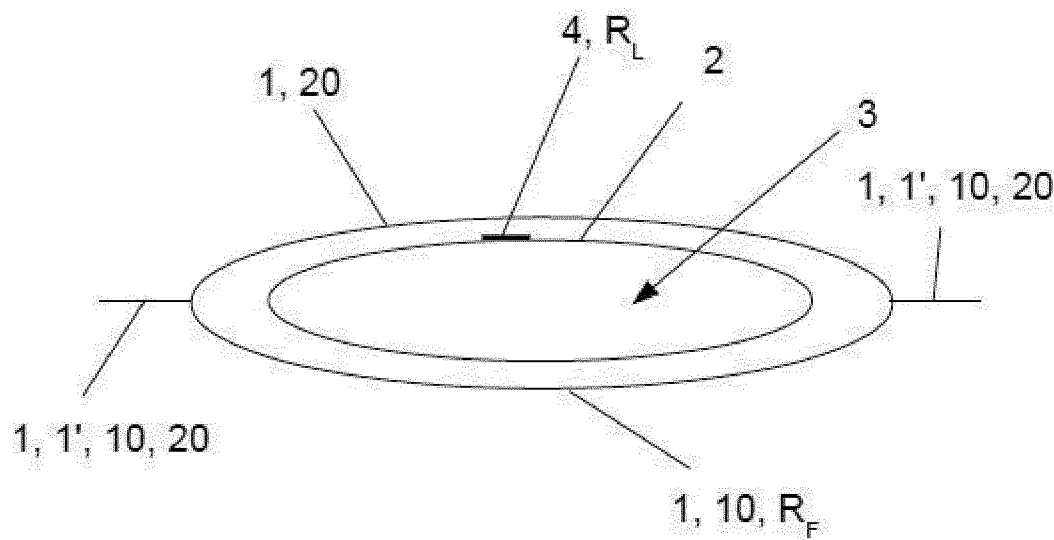


Figure 4:

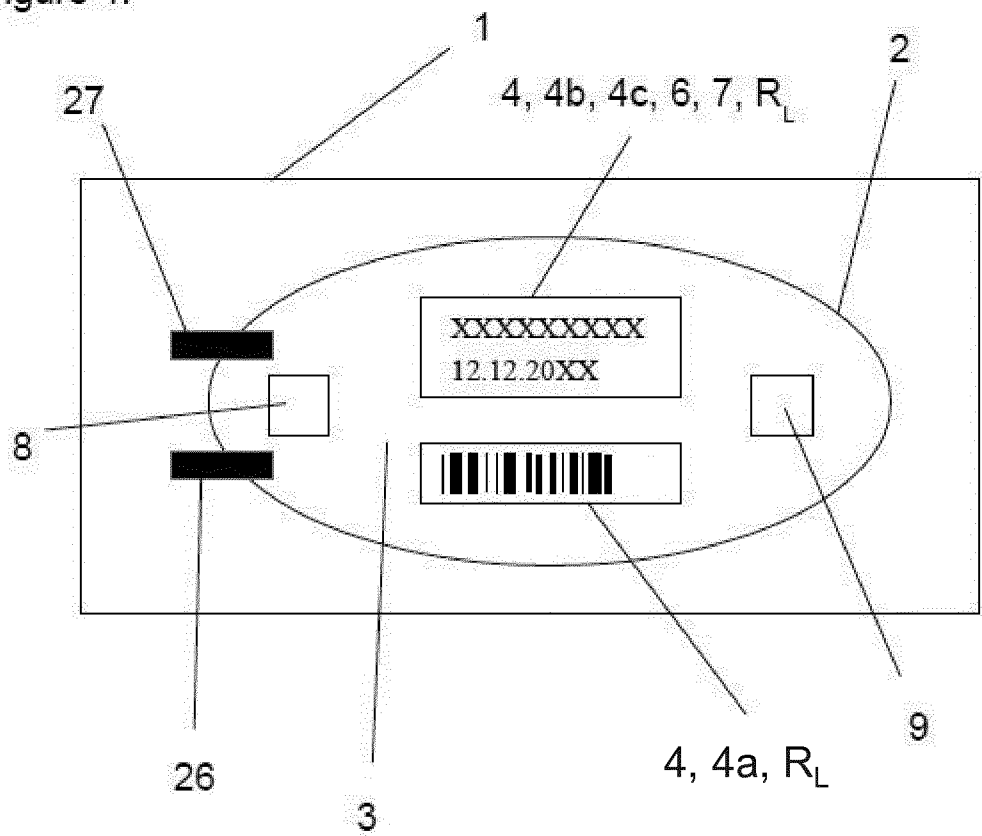


Figure 5

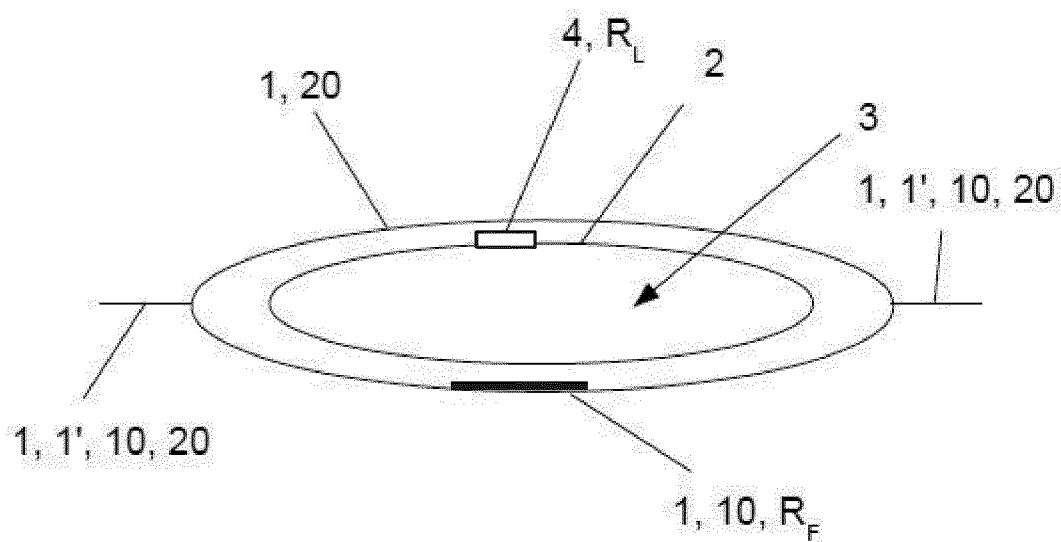
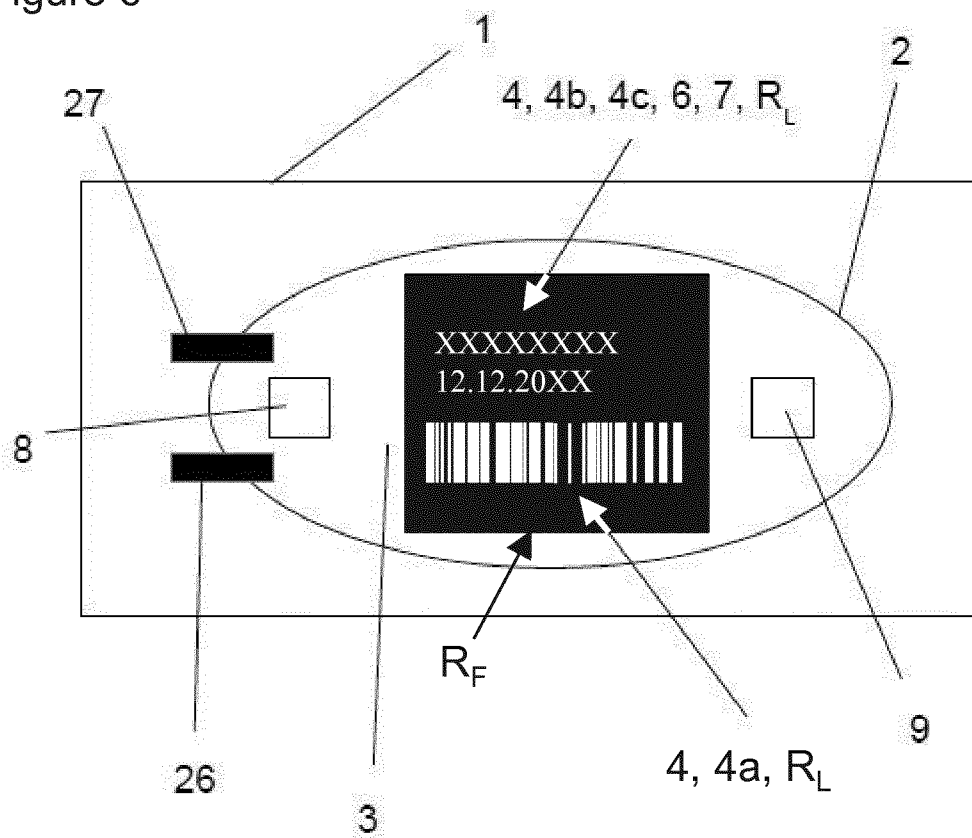


Figure 6



INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2016/051434

A. CLASSIFICATION OF SUBJECT MATTER

INV. A61J1/10 B65D77/04 B65D30/08 B65D33/00
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

B65D A61J

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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Further documents are listed in the continuation of Box C.



See patent family annex.

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Date of the actual completion of the international search

17 February 2016

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

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