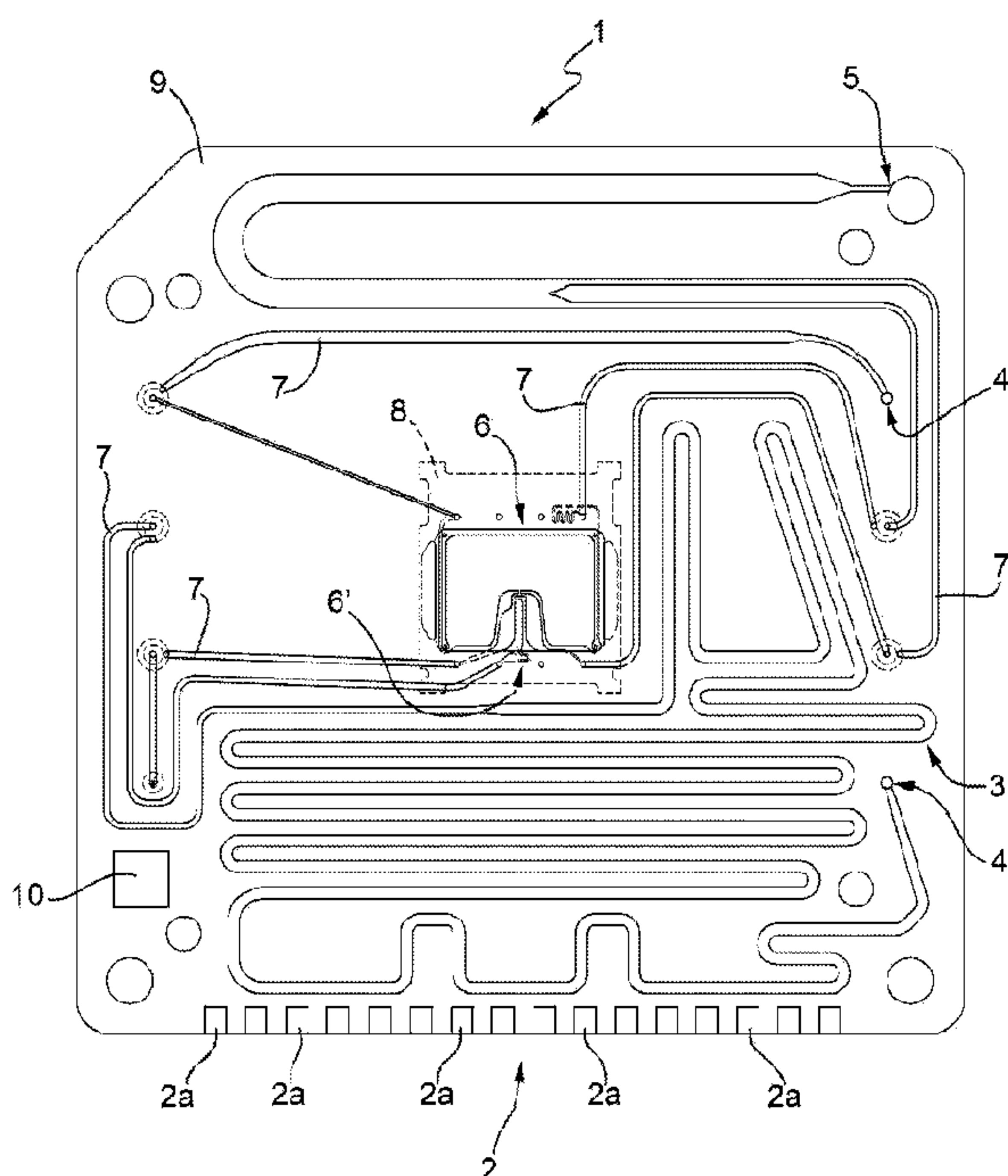




(86) Date de dépôt PCT/PCT Filing Date: 2011/12/22
 (87) Date publication PCT/PCT Publication Date: 2012/06/28
 (45) Date de délivrance/Issue Date: 2019/11/05
 (85) Entrée phase nationale/National Entry: 2014/02/18
 (86) N° demande PCT/PCT Application No.: IB 2011/055920
 (87) N° publication PCT/PCT Publication No.: 2012/085884
 (30) Priorité/Priority: 2010/12/22 (IT BO2010A000755)

(51) Cl.Int./Int.Cl. *B03C 5/02* (2006.01),
B01L 3/00 (2006.01), *G01N 27/447* (2006.01)
 (72) Inventeurs/Inventors:
MEDORO, GIANNI, IT;
MANARESI, NICOLO, IT
 (73) Propriétaire/Owner:
MENARINI SILICON BIOSYSTEMS S.P.A., IT
 (74) Agent: ROBIC

(54) Titre : DISPOSITIF MICROFLUIDIQUE POUR LA MANIPULATION DE PARTICULES
 (54) Title: MICROFLUIDIC DEVICE FOR MANIPULATING PARTICLES



(57) Abrégé/Abstract:

A microfluidic device (1) for isolating particles of at least one given type of a sample; the device (1) is designed to be connected to an apparatus through a plurality of electrical connectors (2a) and comprises a system (3) of microfluidic channels and a flash memory (10), which contains information on the structure (arrangement and geometry of the various components) of the system (3) of microfluidic channels, a map of non-functioning parts of the device (1), the maximum number of uses and the maximum time of use of the device (1); the memory (10) has portions allocated for storing the number of times and the time of use of the device (1).

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

CORRECTED VERSION

(19) World Intellectual Property
Organization
International Bureau



(10) International Publication Number
WO 2012/085884 A8

(43) International Publication Date
28 June 2012 (28.06.2012)

(51) International Patent Classification:

B03C 5/02 (2006.01) *G01N 27/447* (2006.01)
B01L 3/00 (2006.01)

(21) International Application Number:

PCT/IB2011/055920

(22) International Filing Date:

22 December 2011 (22.12.2011)

(25) Filing Language:

Italian

(26) Publication Language:

English

(30) Priority Data:

BO2010A000755 22 December 2010 (22.12.2010) IT

(71) Applicant (for all designated States except US): **SILICON BIOSYSTEMS S.P.A.** [IT/IT]; Via dei Lapidari, 12, Bologna (IT).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **MEDORO, Gianni** [IT/IT]; Via della Badia, 9, I-40033 Casalecchio Di Reno (IT). **MANARESI, Nicolò** [IT/IT]; Via Santo Stefano, 132, I-40125 Bologna (IT).

(74) Agents: **JORIO, Paolo** et al.; STUDIO TORTA S.p.A., Via Viotti, 9, I-10121 Torino (IT).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SC, SD,

[Continued on next page]

(54) Title: MICROFLUIDIC DEVICE FOR MANIPULATING PARTICLES

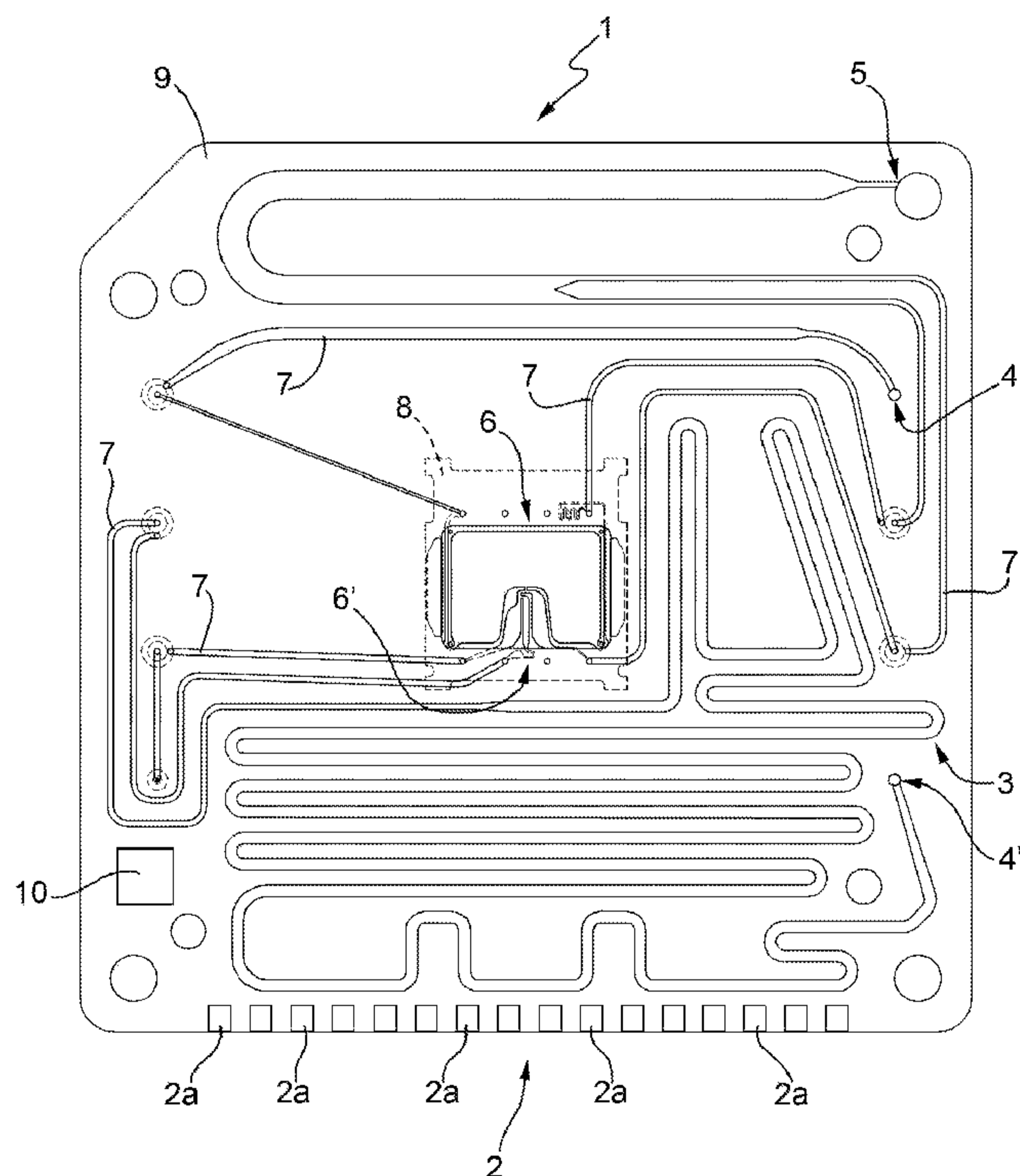


FIG. 1

(57) Abstract: A microfluidic device (1) for isolating particles of at least one given type of a sample; the device (1) is designed to be connected to an apparatus through a plurality of electrical connectors (2a) and comprises a system (3) of microfluidic channels and a flash memory (10), which contains information on the structure (arrangement and geometry of the various components) of the system (3) of microfluidic channels, a map of non-functioning parts of the device (1), the maximum number of uses and the maximum time of use of the device (1); the memory (10) has portions allocated for storing the number of times and the time of use of the device (1).

WO 2012/085884 A8 

SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR,
TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

- (84) Designated States** (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— *with international search report (Art. 21(3))*

(48) Date of publication of this corrected version:

7 September 2012

(15) Information about Correction:

see Notice of 7 September 2012

"MICROFLUIDIC DEVICE FOR MANIPULATING PARTICLES"TECHNICAL FIELD

The present invention relates to a microfluidic device and to
5 uses of such a device.

BACKGROUND OF THE INVENTION

The patent application PCT/IB2010/000615 discloses a
microfluidic device for manipulating particles of a sample (in
10 particular, for isolating a given type of particles from other
particles). Such a device has relatively small dimensions and
can be thrown away after use so as not to contaminate a
possible further sample to be manipulated subsequently. Said
device is typically used in combination with a fixed apparatus
15 of larger dimensions, which comprises an electronic control
unit and governs the various components of the device. The
device is provided with a separation chamber, within which, in
use, manipulation of particles takes place according to what
is described in one or more of the patent applications with
20 publication numbers WO0069565, WO2007010367, WO2007049120, and
WO2007116312.

One and the same apparatus may be able to use different
devices for different uses.

25

Since the devices described above are frequently used in the
diagnostic field, it is very important for them to be managed
with the maximum precision and care. Currently, no system is
envisaged to prevent human-error in management of the devices
30 and/or for driving different components of the devices so as
to optimize operation even in the case of defects of the
devices themselves that are relatively small (and hence
negligible for normal operation thereof).

35 The patent document No. US6726820 describes a microdevice
equipped with an integrated readable, writable, and rewritable

memory.

The patent document No. DE10352887 describes a DNA-Chip-Array-Processor equipped with a memory containing some information.

5

The patent document No. US5690893 describes an analyser equipped with a non-volatile memory containing a condition of analysis.

10 The patent document No. US2004092024 describes a plurality of supporting plates for samples with a resettable memory.

The patent document No. US5384028 describes a biosensor equipped with a memory for storing data that include data
15 regarding fabrication of the biosensor.

The patent document No. WO2005064325 describes a reusable cartridge for bio-analyses including a rewritable non-volatile memory.

20

The patent document No. WO2009137415 relates to droplet actuators.

The patent document No. WO0047322 describes an apparatus for
25 moving elements (packets) along given paths. It does not describe memory media on devices of the disposable type.

None of the documents Nos. US6726820, DE10352887, US5690893, US2004092024, US5384028, WO2005064325, WO2009137415, and
30 WO0047322 envisages a system for driving different components of the devices so as to optimize operation also in the case of defects of the devices themselves that are relatively small (and hence negligible for normal operation thereof).

35 The aim of the present invention is to provide a device and uses of said device that will enable the limits of the prior

art to be overcome, at least partially, and that will, at the same time, be easy and economically advantageous to produce.

SUMMARY

5 According to the present invention, a microfluidic device and uses of said microfluidic device are provided. More particularly, there is provided a microfluidic device for manipulating particles of at least one given type of a sample, the device being designed to be connected to an apparatus and comprising: a connection unit for connecting the device itself
10 to the apparatus, which is external to the device; a microfluidic system, which in turn comprises at least one inlet, at least one outlet, and at least one microfluidic chamber for manipulating the particles of the given type; and a non-volatile rewritable memory containing information on the device chosen
15 in the group consisting of: information on the type of device, information on how the device is to be made to work, at least one identifier of the device, and a combination thereof; the microfluidic chamber comprising a plurality of actuators for moving the particles; the microfluidic device being designed so
20 that the driving of said actuators for moving the particles is carried out by the apparatus; said microfluidic chamber is designed to move selectively the particles of the given type with respect to other particles of the sample; the information on the device comprising an identification of the faulty
25 actuators for moving the particles.

Unless otherwise explicitly specified, in the present text the terms listed below have the meaning indicated hereinafter.

By "equivalent diameter of a section" is meant the diameter of a circle having the same area as the cross section.

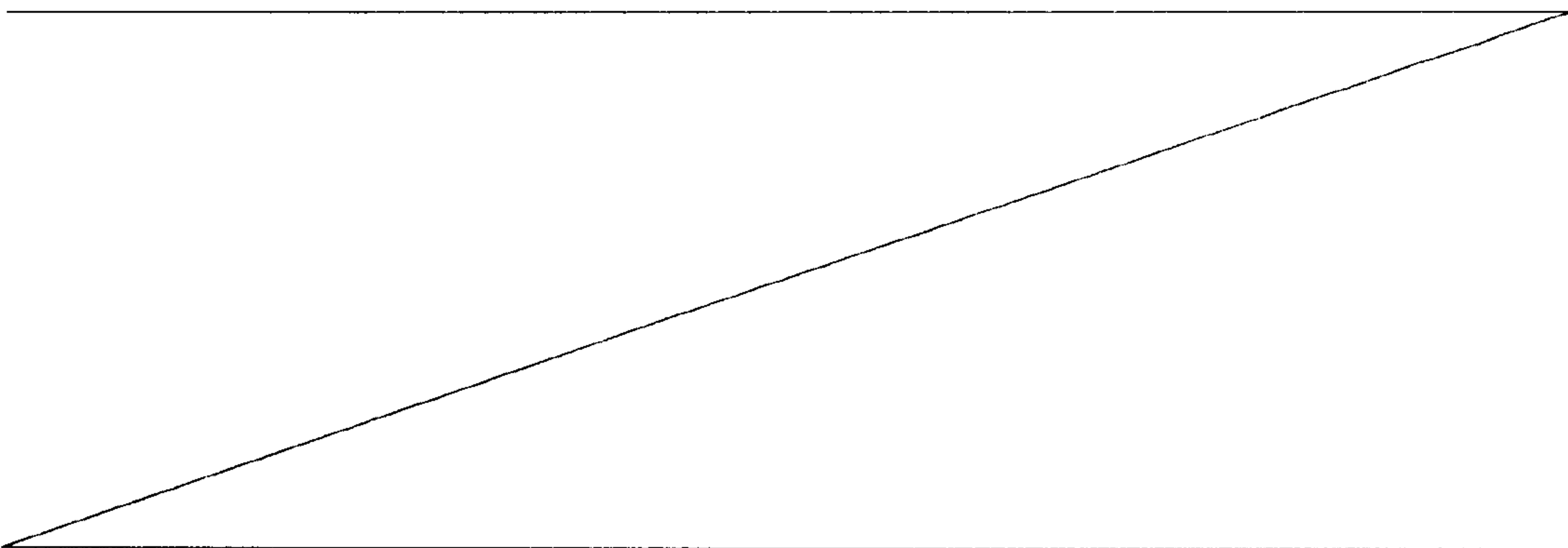
By "section" of a channel or a duct is meant the cross section of the lumen of the channel substantially perpendicular to the longitudinal extension of the channel (or duct), i.e., to the
5 direction of advance of the fluid in the channel (or duct).

By "microfluidic system (or device)" is meant a system (or device) comprising at least one microfluidic channel (or duct) and/or a microfluidic chamber.

10 By "microfluidic chamber" is meant a chamber delimiting an internal space having one of its own dimensions (in particular, the height) of less than 1 mm.

By "microfluidic channel (or duct)" is meant a channel (or duct) having a cross section with equivalent diameter of less
15 than 1 mm.

The dimensions of the channels or ducts or chambers can be measured in a standard way with profilometers.



In the present text, by "particle" is meant a corpuscle having the largest dimension of less than 500 μm (advantageously, less than 150 μm). Non-limiting examples of particles are: cells, cellular detritus (in particular, cellular fragments),
5 cellular aggregates (such as, for example, small clusters of cells deriving from stem cells such as neurospheres or mammospheres), bacteria, lipospheres, (polystyrene and/or magnetic) microspheres, nanospheres (for example, nanospheres of up to 100 nm), complexes formed by microspheres bound to
10 cells, etc. Advantageously, the particles are cells.

According to some embodiments, the particles (advantageously cells and/or cellular detritus) have the largest dimension of less than 60 μm .

15

The dimensions of the particles can be measured in a standard way with microscopes with graduated scale or normal microscopes used with slides (deposited on which are the particles) with graduated scale.

20

In the present text, by "dimensions of a particle" is meant the length, width, and thickness of the particle.

25

The term "substantially selective" is used for identifying a displacement (or other similar terms indicating a movement and/or a separation) of particles, where the particles that are displaced and/or separated are particles that for the most part are constituted by one or more given types. Advantageously, a substantially selective displacement (or
30 other similar terms indicating a movement and/or a separation) envisages displacing particles with at least 90% (advantageously 95%) of particles of the given type or types (percentage given by the number of particles of the given type/ types with respect to the total number of particles).

35

BRIEF DESCRIPTION OF THE DRAWING

The invention is described hereinafter with reference to the annexed plate of drawings, which illustrates a non-limiting example of embodiment thereof, and in which:

Figure 1 is a schematic illustration of a system built in accordance with the present invention

EMBODIMENTS OF THE INVENTION

In accordance with a first aspect of the present invention, a microfluidic device is provided for manipulating at least one given type of particles of a sample. In particular, the device 1 is designed to isolate the aforementioned particles with respect to further particles in a substantially selective way. More in particular, the device is of the disposable type.

In Figure 1, designated as a whole by 1 is one such device.

The device 1 is designed to be connected to an apparatus (not illustrated) and comprises a connection unit 2 for connecting the device 1 itself to the apparatus electrically (or electromagnetically).

The apparatus is designed to drive different components of the device 1. The apparatus is external to (separate from) the device 1 and can come into contact therewith via the connection unit 2. According to some embodiments, the apparatus has a seat (not illustrated), in which, in use, the device 1 is inserted so as to connect the apparatus to the device 1.

According to the depicted embodiment, the connection unit 2 comprises a plurality of electrical connectors 2a (of which only some are schematically illustrated in Figure 1), each of which is designed to couple with corresponding connectors (not illustrated) of the apparatus. Advantageously, the connectors 2a are at least partially exposed towards the outside of the device 1. In use, through the connectors 2a the device 1

receives operating commands from the apparatus.

According to further not shown embodiments, the connection unit 2 is able to communicate (receive and/or transmit information and/or commands) in wireless mode (in particular, with the apparatus). This can be particularly useful when it is desired to identify one or more given devices in a group of devices.

By means of the wireless communication it is possible to prevent direct contact between the device 1 and the apparatus. This presents a dual advantage: (i) in the case where the device 1 has been used (biological waste) it prevents contamination of the apparatus; (ii) in the case the device 1 is new, it prevents contamination of the device 1 itself (possibly sterile or DNA/RNA-free).

The device 1 comprises a microfluidic system 3, which in turn has at least one inlet 4, at least one outlet 5, and at least one (microfluidic) chamber 6, within which, in use, the particles of the given type are manipulated. In particular, the system 3 also comprises at least one microfluidic channel 7. More in particular, the system 3 comprises a plurality of microfluidic channels 7.

According to the embodiment depicted in Figure 1, the system 3 comprises two inlets 4 and 4', two outlets 5 and 5', and two (microfluidic) chambers 6 and 6'. In the specific case, the chamber 6' comprises three portions of a different shape connected to one another.

The chamber 6 is designed to move the particles of the given type selectively with respect to the further particles of the sample. In particular, in use, the sample is brought into the chamber 6, and the particles of the given type are selectively brought into the chamber 6' previously filled with a liquid

transport medium. The particles of the given type are then recovered by an operator through the outlet 5'.

The device 1 comprises a substructure 8, set in a position corresponding to which is the chamber 6 (and the chamber 6'). More in particular, the substructure 8 delimits at the bottom the chamber 6 (and the chamber 6'). The substructure 8 is mainly (more in particular, completely) made of silicon.

10 The device 1 comprises a plurality of actuators (for example, valves and/or electrodes), which are operated on the basis of commands coming from the apparatus. According to some embodiments, the chamber 6 comprises a plurality of actuators (for moving the particles). The actuators of the device 1
15 comprise the actuators of the chamber 6 (and/or 6'). The actuators of the chamber 6 (and/or 6') comprise (are), in particular, electrodes.

Driving of the various actuators (electrodes) of the chamber 6
20 enables displacement of the particles (and is governed by the apparatus). Advantageously, the structure and operation of the chamber 6 (and of the chamber 6') is in accordance with what is described in one or more of the patent applications with publication numbers WO0069565, WO2007010367, WO2007049120 and
25 WO2007116312 filed in the name of the present applicant.

More in particular, the device 1 presents the same structure and operation as the device described in the patent application PCT/IB2010/000615 (publication No. WO2010106434)
30 filed in the name of the present applicant. The device 1 differs from the one described in the patent application PCT/IB2010/000615 owing to the presence of a partially rewritable memory as described hereinafter. Likewise, the apparatus presents the same structure and operation as the
35 apparatus described in the patent application PCT/IB2010/000615.

According to the depicted embodiment, the device 1 also comprises an element 9 (in particular, made of plastic material), made in which are the inlet 4 (and 4') and the outlet 5 (and 5'). Also made in the element 9 are the microfluidic channels 7 and (partially) the chamber 6 (and 6'). More in particular, the system 3 is at least partially (according to some embodiments, completely) made in the element 9.

10

Advantageously, the element 9 comprises (more specifically, mainly consists of) plastic material.

15

According to specific embodiments, the device 1 comprises a printed circuit board (PCB - not shown). The PCB is connected to the connection unit 2 and to the substructure 8. The PCB is set underneath the element 9.

20

The device 1 further comprises a non-volatile rewritable memory (a flash memory). More in particular, the device 1 comprises a physical medium 10 for said memory.

25

According to some embodiments (as the one depicted), the physical medium 10 of the memory is set in the area of the element 9 separately (i.e., in a different position) from the substructure 8. In this way, it is possible to reduce the manufacturing costs.

30

According to further embodiments (not illustrated), the physical medium 10 of the memory is set in a position corresponding to the substructure 8. The physical medium 10 is connected to the PCB.

35

In particular, the memory contains information on the device 1.

Advantageously, the memory contains information on the type of device 1. Advantageously, the information on the type of device 1 is stored at least partially (in particular, completely) in a non-rewritable (read-only) area of the memory.

In particular, the information on the type of device 1 comprises information on the structure (arrangement and geometry of the various components) of the device 1. More in particular, the information on the type of device 1 comprises information on the structure (arrangement and geometry of the various components) of the microfluidic system 3. In this way, the apparatus (more in particular, the control unit of the apparatus) is able to recognize the device 1 and drive its own actuators (and hence the various components of the device 1 itself) in an appropriate way. The information on the structure of the device 1 (in particular, of the microfluidic system 3) is written in the memory at the time of testing of the device 1 and is of the read-only type (i.e., it cannot be modified).

According to some embodiments, the information on the type of device 1 comprises information on the structure (arrangement and geometry of the various components) of the PCB. In this way, the apparatus (more in particular, the control unit of the apparatus) is able to recognize the device 1 and drive the PCB in an appropriate way. The information on the structure of the PCB is written in the memory at the time of testing of the device 1 and is of the read-only type (i.e., it cannot be modified).

Advantageously, the information on the type of device 1 comprises an indication of the actuators (in particular, electrodes) of the device 1 that are faulty. In particular, the indication of the faulty actuators comprises (more in particular, is) the indication of the faulty actuators (in

particular, electrodes) of the chamber 6 (and/or 6').

More precisely, the information on the type of device 1 comprises the position/ positions of the faulty electrode/electrodes. In other words, the memory comprises a mapping of the faulty electrodes. This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be modified). Typically, the faulty electrodes are non-
10 functioning or in any case unable to guarantee proper operation.

With this type of information, the apparatus (in particular, the control unit of the apparatus) manipulates (more
15 precisely, moves) the particles (in particular, the particles of the given type) so as to prevent area/areas where the faulty electrode/electrodes is/are set. In other words, the electrodes are actuated so as to impose on the particles one or more paths that do not pass in a position corresponding to
20 the faulty electrode/electrodes.

In this way, the very important advantage is achieved of rendering possible use of the device 1 even when it has small defects.

25

This enables significant reduction in the number of rejects of the devices 1 at the end of production, when the devices 1 are tested, without reducing the quality of the results in use of the devices 1 themselves. It is thus possible to reduce
30 considerably the manufacturing costs.

According to some embodiments, the information on the type of device 1 comprises the maximum number of uses that can be performed. This information is written in the memory at the
35 time of testing of the device 1. This information is of the read-only type (i.e., it cannot be modified).

Advantageously, the memory is designed to store the number of times that the device 1 is used. In particular, a portion of the memory is allocated for storage of the number of times
5 that the device 1 is used.

In use, whenever the device 1 is used, the apparatus (in particular, the control unit of the apparatus) records said use in the memory. When the apparatus (in particular, the
10 control unit of the apparatus) detects (by reading the memory) that the maximum number of uses has been reached, the apparatus itself blocks operation and issues a corresponding error signal.

15 In this way, it is possible to guarantee a high quality level in use of the device 1. In particular, for example, in its use for diagnostic purposes, the device 1 can be used just once to prevent contamination of the samples. In this case, what is described enables prevention of errors by the operator and,
20 hence, potential incorrect results.

According to some embodiments, the information on the type of device 1 contains the maximum time of use of the device 1. This information is written in the memory at the time of
25 testing of the device 1. This information is of the read-only type (i.e., it cannot be modified).

Advantageously, the memory is designed to store the time during which the device 1 is used. In particular, a portion of
30 the memory is allocated for storage of the time during which the device 1 is used.

In use, whenever the device 1 is used, the apparatus (in particular, the control unit of the apparatus) records the
35 time of said use in the memory and possibly adds it to the time already present in the memory itself. When the apparatus

(in particular, the control unit of the apparatus) detects (by reading the memory) that the maximum time of use has been reached, the apparatus itself blocks operation and issues a corresponding error signal.

5

In this way, it is possible to guarantee a high quality level in use of the device 1. In particular, it is thus possible to prevent a device 1 that is worn out (owing to excessive use) from being used again.

10

Advantageously, the memory is accessible via password. In this way, it is possible to reduce the risks of said memory being tampered with.

15

Advantageously, the memory is encrypted. In this way, the possibility of the device 1 being used in combination with unsuitable apparatuses is prevented.

20

Advantageously, the information on the type of device 1 comprises the expiry date of the device 1. This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be modified). In use, when the apparatus (in particular, the control unit of the apparatus) detects (by reading the memory) that the expiry date is passed, the apparatus itself blocks operation and issues a corresponding error signal.

25

Also in this way the quality and safety of operation is improved.

30

According to some embodiments, the information on the type of device 1 comprises the position of the microfluidic system 3 (in the case where the latter has been produced by means of lithographic techniques) with respect to the substructure (wafer) 8. The memory further comprises the identifier of the substructure (wafer) 8 within each production lot and the

35

identifier of the lot itself. This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be modified).

5

According to some embodiments, the information on the type of device 1 comprises an indication on whether the information on the faulty actuators (electrodes) is present or not (for example: 1 = yes, 0 = no). This indication is written in the

10 memory at the time of testing of the device 1 and is of the read-only type (i.e., it cannot be modified).

According to some embodiments, the information on the type of device 1 comprises an indication on the type of error (for

15 example, a code and/or a coloured marker). Advantageously, the information on the type of device 1 moreover contains an indication of the model of the device 1 (each revision or new version of the device 1 can require different operating procedures that can possibly also be stored in the memory).

20

According to some particular embodiments, the information on the type of device 1 contains an indication on the type of mask (or masks) that has (have) been used for producing the element 9.

25

According to some embodiments, the information on the type of device 1 comprises information on the bonding system that has been used for producing the device 1. This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be

30 modified).

35

According to some embodiments, the information on the type of device 1 comprises information on the height of the chamber 6 (and/or 6') (for example, 90 μm). This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be

modified).

According to some embodiments, the information on the type of device 1 comprises information on the maximum number of droplets that can be recovered per sample (in particular through the 'outlet 5'). Said droplets are the ones that contain (or in any case should contain) the particles of the given type. This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be modified).

According to some embodiments, the information on the type of device 1 comprises information on the average volume of the aforementioned droplets. This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be modified).

According to some embodiments, the information on the type of device 1 comprises information on the authorisation of the device 1 for use of different transport solutions (for example PBS - phosphate buffered saline solution). This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be modified).

In use, the transport solution is brought into the chamber 6' prior to selective displacement of the particles of the given type.

According to some embodiments, the information on the type of device 1 comprises information on the range of operating temperatures (for example, 277K-309K). This information is written in the memory at the time of testing of the device 1. This information is of the read-only type (i.e., it cannot be modified).

According to some embodiments, the memory is designed to contain (in particular, has a portion allocated for storing) information on whether the device 1 has been inserted in the apparatus but not expelled (for example: 1 = yes, 0 = no).
5 This information is written in the memory at the time of use of the device 1 (by the apparatus - more in particular, by the control unit of the apparatus). According to some embodiments, upon charging (in particular, immediately after charging), the
10 information on charging is written (for example by writing 1). Upon expulsion (in particular, immediately prior to expulsion), the information on the expulsion is written (for example, by writing 0).

15 This information indicates when the device 1 has been charged into the apparatus but has never been expelled therefrom and has the purpose of enabling prosecution of the experiment in the case of reboot of the apparatus subsequent to charging. When the device 1 is removed with the apparatus turned off
20 manually by a technician, the device 1 should no longer be used, and it is advantageous for the apparatus to be able to detect when the memory has this variable equal to 0 and consequently understands that the device 1 has been moved away and may have been tampered with, thus preventing
25 continuation of the experiment.

According to some embodiments, the memory is designed to contain (in particular, has a portion allocated for storing) one or more of the following items of information: date and
30 time of start of experiment (writing in the memory is performed at start of the experiment); date and time of insertion of the device 1 into the apparatus 1 (writing in the memory is performed immediately after insertion); date and time of the end of the experiment (writing in the memory is
35 performed at the end of the experiment); date and time of expulsion of the device 1 from the apparatus (writing in the

memory is performed just before expulsion); set of parameters used (in particular, transport solutions used - for example, PBS) (writing in the memory is performed at start of the experiment); an identifier of the apparatus used (writing in
5 the memory is performed immediately after insertion); name of the pattern used (i.e., the configuration of the electrophoresis field, i.e., the association between each electrode and each phase - V_p or else V_m (the energizing potentials that can be assumed by the electrodes)) (writing in
10 the memory is performed at start of the experiment); frequency, amplitude and phase delta for V_p , V_m and V_{lid} (the potential for energizing the lid) (writing in the memory is performed at start of the experiment).

15 According to some embodiments, the information on the type of device 1 comprises a combination of a number of the types of information referred to above.

Advantageously, the memory contains information on how the
20 device 1 must be made to work (namely, information on which operating procedures must be implemented, for example, how the various actuators are to be actuated).

According to some variants, the information contains
25 indications of the use (for example, for research use only or else for *in vitro* diagnostics).

According to some embodiments, the memory contains information chosen from among: information on the type of device 1,
30 information on how the device 1 is to be made to work, and a combination thereof.

Advantageously, the information on how the device 1 is to be made to work is stored at least partially (in particular,
35 completely) in a non-rewritable (read-only) area of the memory.

According to some embodiments, the memory comprises at least one identifier of the device 1 (namely, a unique identifier code of the device. 1). In other words, the memory of each device 1 contains an identifier different from the identifier of the other devices. This information is written in the memory at the time of testing of the device 1.

Advantageously, the identifier of the device 1 is stored at least partially (in particular, completely) in a non-rewritable (read-only) area of the memory.

According to some embodiments, the memory contains information chosen from among: information on the type of device 1, information on how the device 1 is to be made to work, at least one identifier of the device 1, and a combination thereof.

Advantageously, one or more of the items of information present in the memory can be displayed by means of a human-machine interface (HMI - for example a screen) of the apparatus.

According to a second aspect of the present invention, use of the device 1 for diagnostic purposes is provided.

According to a third aspect of the present invention, use of the device 1 for separating the particles of the given type from other particles is provided.

Unless otherwise explicitly indicated, the contents of the references (articles, books, patent applications, etc.) cited in this text are integrally recalled herein. In particular, the aforementioned references are herein incorporated by reference.

CLAIMS

- 1.- A microfluidic device for manipulating particles of at least one given type of a sample, the device (1) being designed to be
5 connected to an apparatus and comprising: a connection unit (2) for connecting the device (1) itself to the apparatus, which is external to the device (1); a microfluidic system (3), which in turn comprises at least one inlet (4; 4'), at least one outlet (5; 5'), and at least one microfluidic chamber (6; 6') for
10 manipulating the particles of the given type; and a non-volatile rewritable memory containing information on the device (1) chosen in the group consisting of: information on the type of device (1), information on how the device (1) is to be made to work, at least one identifier of the device (1), and a
15 combination thereof; the microfluidic chamber comprising a plurality of actuators for moving the particles; the microfluidic device (1) being designed so that the driving of said actuators for moving the particles is carried out by the apparatus; said microfluidic chamber is designed to move
20 selectively the particles of the given type with respect to other particles of the sample; the information on the device (1) comprising an identification of the faulty actuators for moving the particles.
- 25 2.- The device according to claim 1, wherein the information on the device (1) is at least partially set in a non-rewritable area of the memory.
- 30 3.- The device according to claim 1 or 2, wherein the actuators are operated on the basis of commands coming from the apparatus.

- 4.- The device according to any one of claims 1 to 3, wherein the actuators of the chamber comprise electrodes.
- 5 5.- The device according to any one of claims 1 to 4, wherein the information on the device (1) comprises information on the structure of the microfluidic system (3).
- 10 6.- The device according to claim 5, wherein the information on the structure of the microfluidic system (3) comprises information on arrangement and geometry of different components of the microfluidic system (3).
- 15 7.- The device according to any one of claims 1 to 6, wherein the information on the device (1) comprises the maximum number of uses that are available for being performed.
- 20 8.- The device according to any one of claims 1 to 7, wherein the memory is designed to store the number of times that the device (1) is used.
- 25 9.- The device according to claim 8, wherein a portion of the memory is allocated for storage of the number of times that the device (1) is used.
- 30 10.- The device according to any one of claims 1 to 9, wherein the information on the device (1) comprises the maximum time of use of the device (1).
- 11.- The device according to any one of claims 1 to 10, wherein

the memory is designed to store the time during which the device (1) is used.

5 12.- The device according to claim 11, wherein a portion of the memory is allocated for storage of the time during which the device (1) is used.

10 13.- The device according to any one of claims 1 to 12, wherein the connection unit (2) comprises electrical connectors (2a) at least partially exposed towards the outside of the device (1).

14.- The device according to any one of claims 1 to 13, wherein the memory is accessible by means of a password.

15 15.- The device according to any one of claims 1 to 14, wherein the memory is encrypted.

20 16.- The device according to any one of claims 1 to 15, and comprising: a silicon substructure (8), in the area of which the chamber (6; 6') is located; an element (9) mainly made of plastic material, in which said inlet (4; 4'), said outlet (5; 5'), and at least one channel (7) of the microfluidic system (3) are obtained; and a physical medium (10) for said memory, said physical medium (10) being set in an area corresponding to
25 the element (9) mainly made of plastic material in a position different from the silicon substructure (8).

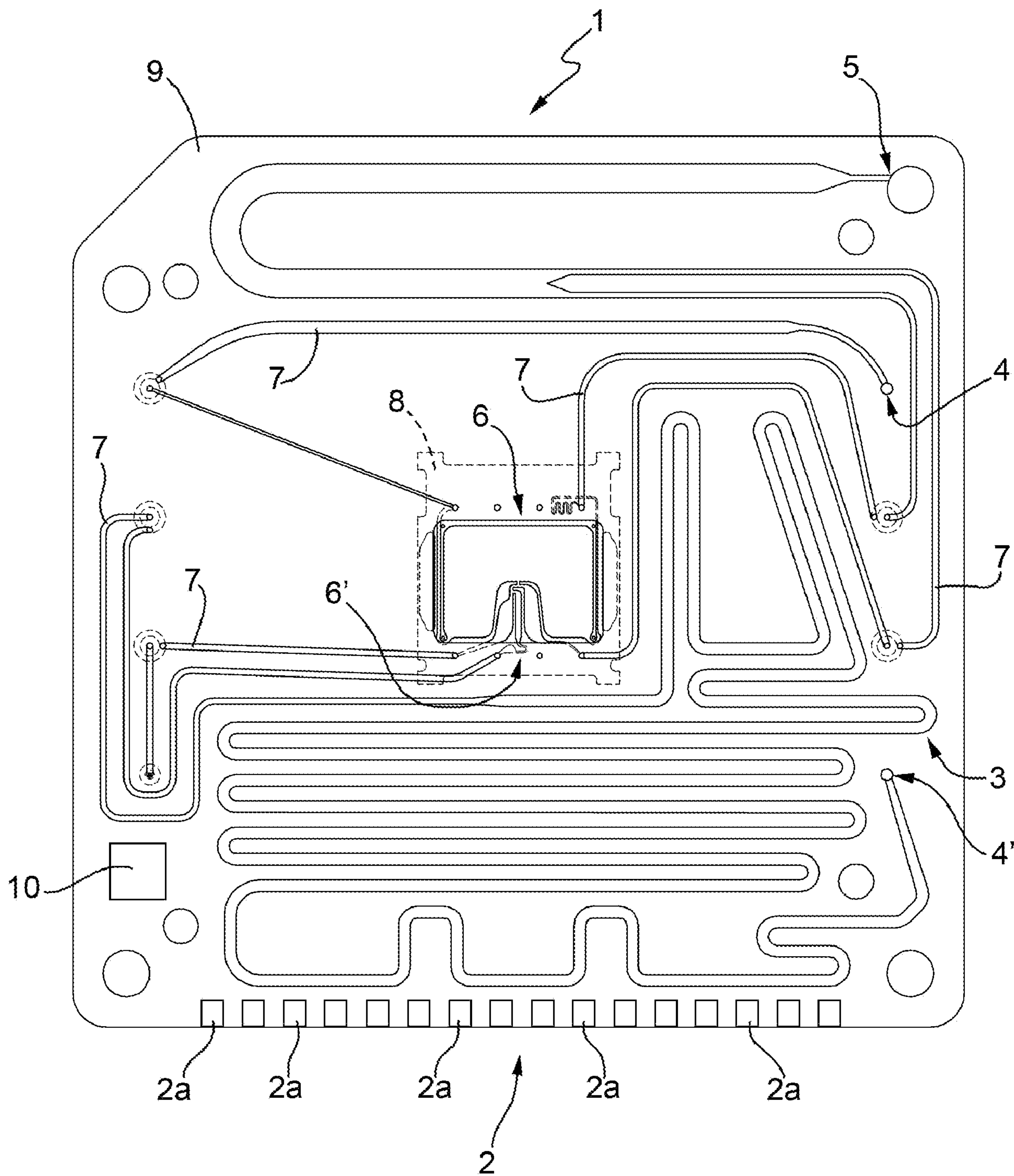


FIG. 1

