



- (51) **International Patent Classification:**
C12M 1/00 (2006.01) *C12M 1/26* (2006.01)
- (21) **International Application Number:**
PCT/US2016/037154
- (22) **International Filing Date:**
13 June 2016 (13.06.2016)
- (25) **Filing Language:** English
- (26) **Publication Language:** English
- (30) **Priority Data:**
14/736,623 11 June 2015 (11.06.2015) US
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(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, BN, 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, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, 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, ST, SZ,

[Continued on next page]

(54) **Title:** ENHANCED PHOTOBIOREACTOR SYSTEM

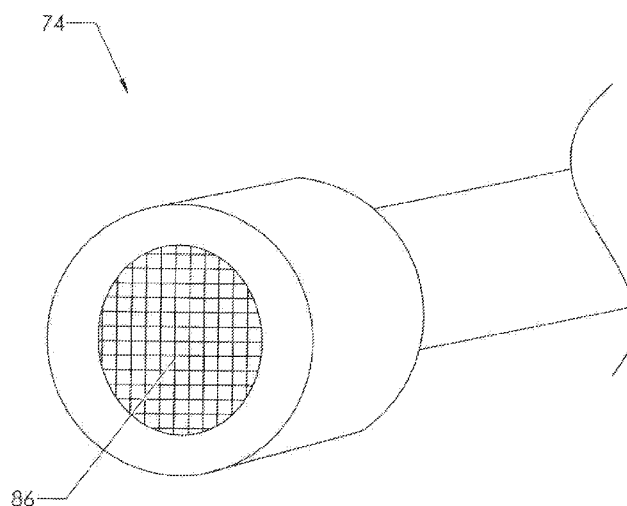


FIG. 7

(57) **Abstract:** A space efficient photo-bioreactor. The bioreactor grows microalgae in a tall array of transparent flooded tubes. A nutrient media is circulated through the tubes. The array is configured to maximize the amount of sunlight tailing upon each tube so that growth of the microalgae is as uniform as possible. A vertically-oriented gasser tube is provided. Gas is injected into this gasser tube along with the liquid nutrient medium. A bubble-size limiter is employed in the gas injector. The flow rates are configured so that the liquid nutrient medium and injected gas remain within the vertical gasser tube for 30 seconds or more.





TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, 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, KM, ML, MR, NE, SN, TD, TG).

Published:

- *with international search report (Art. 21(3))*
- *with information concerning one or more priority claims considered void (Rule 26bis.2(d))*

Declarations under Rule 4.17:

- *as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))*

TITLE OF THE INVENTION:
ENHANCED PHOTOBIOREACTOR SYSTEM

CROSS-REFERENCES TO RELATED APPLICATIONS

This non-provisional patent application is a continuation-in-part of prior U.S. application serial number 13/271,622, which itself claims the benefit, pursuant to 37 C.F.R. section 1.53(c), of an earlier-filed provisional patent application. The earlier application was assigned serial number 61/392,053. It listed the same inventor.

**STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR
DEVELOPMENT**

Not Applicable.

MICROFICHE APPENDIX

Not Applicable member

DESCRIPTION

Title of the Invention: Enhanced Photobioreactor System

1. Technical Field.

5 This invention relates to the field of renewable energy. More specifically, the invention comprises a space-efficient photo-bioreactor and methods for controlling the bioreactor.

2. Background Art.

10 The continued use of petroleum-derived fuels is now widely seen as unsustainable. However, much of the existing transportation structure is dependent upon the combustion of liquid fuels. Changing to a completely different energy source – such as battery power – is at present unrealistically expensive and inefficient.

On the other hand, presently available biofuels can be substituted for petroleum-derived fuels without the need for extensively modifying existing internal combustion engines. One promising alternative fuel is biodiesel, which can be substituted for petroleum
15 diesel in many modern engines (albeit with a slight reduction in specific energy).

Oil crops can be used to make biodiesel. These are attractive, as the total cycle of production through consumption can be made carbon-neutral. Unfortunately, though, oil crops are not very space-efficient. It is estimated that if 24% of the total cropland in the United States was devoted to a high-yielding oil crop such as palm oil, this would still only
20 meet about half of the demand for transportation fuels.

Microalgae-based bio-fuels hold the promise of much greater space efficiency. Like plants, microalgae use sunlight to produce oils. They do it much more efficiently than crop plants, though. Microalgae-based biodiesel is still in a developmental state in terms of cost efficiency. However, it is clear that biodiesel can be made from microalgae. In order to
25 make such a process economically efficient, it is important to use as many of the products produced as possible.

It is known to cultivate algae in a series of ponds. While this method does work, it is not space efficient. In fact, using the open-pond method, more surface area is needed to grow a given mass of biofuel than would be needed for conventional row crops yielding the same
30 amount of biofuel. In order to realize the potential of microalgae-based fuels, then, it is far preferable to provide a more space-efficient system.

Further, prior art photo-bioreactors typically require concentrated carbon dioxide as a feed material for the photosynthesis. Concentrated carbon dioxide must be collected, transported, and stored. Even if an on-site source is available, it must still be segregated from the surrounding air. It would be preferable to provide a photo-bioreactor that can run on ordinary air. Such a reactor would still be useful in removing carbon dioxide from the atmosphere, but would eliminate the need to separately collect and store the carbon dioxide. The present invention is able to run on ordinary air. The present invention is also quite space efficient. These and other advantages will be explained in the following descriptions.

SUMMARY OF INVENTION

The present invention comprises a space efficient photo-bioreactor system. The bioreactor grows microalgae in a tall array of transparent flooded tubes. A nutrient media is circulated through the tubes. The array is configured to maximize the amount of sunlight falling upon each tube so that growth of the microalgae is as uniform as possible.

In the preferred embodiments a vertical support structure is provided for the array of tubes. A reservoir is located on the top of this structure. Flow from the reservoir branches into multiple, independent flow paths. Each independent flow path includes a serpentine array of transparent tubes. A liquid pump or pumps collects the flow from the flow paths, pressurizes it, and pumps it back to the reservoir. A vertically-oriented gassing tube carries the flow from the pump back up to the reservoir. Air, or other CO_2 -containing gas, is injected near the bottom of the gassing tube. The size of the gas bubbles is controlled by injecting through appropriate metering openings. The gas diffuses through the liquid medium as the mixture rises in the gasser tube.

Microalgae are harvested from the photo-bioreactor and processed for various suitable uses. One use is the manufacturing of biodiesel. The microalgae is filtered and dried. Lipids are then extracted from the microalgae. These lipids are made into biodiesel through a transesterification process. The lipids may be used to make other products as well.

Some of the biodiesel can be used to run a diesel engine to furnish electrical and/or mechanical power to the bioreactor. Exhaust gas emitted by the diesel engine is preferably fed back into the bioreactor. Carbon dioxide from other greenhouse gas sources is preferably also fed into the bioreactor.

65 BRIEF DESCRIPTION OF DRAWINGS

FIG. 1 is a schematic view, showing the operation of the photo-bioreactor and other related processes.

FIG. 2 is an elevation view showing the arrangement of the bioreactor tubes.

70 FIG. 3 is a perspective view, showing a typical circulation path for the bioreactor tubes.

FIG. 4 is an exploded perspective view, showing a typical gassing/degassing system.

FIG. 5A is a side elevation view, showing an exemplary photo-bioreactor.

FIG. 5B is a perspective view, showing the photo-bioreactor of FIG. 5A.

FIG. 6A is a front elevation view, showing the photo-bioreactor of FIG. 5A.

75 FIG. 6B is a front elevation view, showing the photo-bioreactor of FIG. 5A.

FIG. 7 is a detailed perspective view, showing a mesh that can be used in the air injector to limit bubble size.

REFERENCE NUMERALS IN THE DRAWINGS

80	10	energy harvesting system	12	water tank
	14	nutrients	16	nutrient tank
	18	photo-bioreactor	20	harvesting unit
	22	filtering unit	24	drying unit
	26	lipids extraction unit	28	trans-esterification unit
85	30	biodiesel	32	diesel engine
	34	carbon dioxide input	36	inoculum input
	38	support frame	40	rack
	42	bioreactor tube	43	sunlight
	44	elbow	46	gassing/degassing system
90	48	housing	50	carbon dioxide inlet
	52	oxygen outlet	54	aluminum helix
	56	coolant inlet	58	coolant outlet
	60	inlet	62	outlet
	64	inlet manifold	66	outlet manifold
95	68	reservoir	70	liquid pump

	72	gasser tube	74	gas injector
	76	gas injector pump	78	gas inlet
	80	tube column	82	tube column
	84	tube row	86	mesh
100	88	pump intake line	90	pump discharge line
	92	reservoir outlet line		

DESCRIPTION OF EMBODIMENTS

FIG. 1 shows a schematic view of a comprehensive energy harvesting system 10 based on one or more photo-bioreactors 18. The photo-bioreactors are preferably made as vertical structures having a relatively small “footprint” compared to the volume of liquid media they contain.

Nutrients 14 are mixed with water from water tank 12 (or other suitable water source) to create a nutrient medium which is preferably stored in nutrient tank 16. Inoculum input 36 is fed into a portion of the nutrient medium and this mixture is then fed into the photo-bioreactors.

Sunlight falling on the photo-bioreactors causes microalgae to grow inside. This is eventually harvested in harvesting unit 20. The product of the harvesting unit is then fed through filtering unit 22, where the microalgae is removed and residual nutrient medium is sent back to the photo-bioreactors.

The microalgae is then fed from filtering unit 22 to drying unit 24, where it is dried. The dried microalgae is then fed through lipids extraction unit 26. The extracted lipids are then sent to trans-esterification unit 28, which converts the lipids to biodiesel 30 using processes well known to those skilled in the art. The “waste” products from the lipids extraction unit are preferably fed back to the bioreactors.

The biodiesel thus produced can be transported and used as a substitute for conventional fuels. A portion of the biodiesel produced can also be used to run an on-site diesel generator. The generator can then provide power for the energy harvesting system 10.

The system preferably re-uses the products of each stage in the process. For example, the carbon dioxide produced by the on-site generator is preferably fed back into the bioreactors. More carbon dioxide will likely be needed for this embodiment – and this is furnished via carbon dioxide input 34.

FIG. 2 shows a partial sectional elevation view through one of the photo-bioreactors. As mentioned previously, each photo-bioreactor preferably has a small footprint in comparison to the volume it contains. Support frame 38 supports a number of layered racks 40. Each rack 40 supports a number of bioreactor tubes 42. The tubes are relatively thin-walled transparent structures oriented perpendicularly to the view in FIG. 2. They are spaced (both horizontally and vertically) so that sunlight 43 can pass into the bioreactor and fall on each of the tubes.

The liquid nutrient medium flows through the tubes. The tubes are joined together so that an elongated flow path is created. FIG. 3 shows one approach to joining the tubes in one rack 40. Each tube has an inlet end and an outlet end. The terms "inlet end" and "outlet end" are arbitrary terms depending on the flow direction through a particular tube. Two adjacent tubes may be joined by installing an elbow 44 between the outlet end of one tube and the inlet end of the adjacent tube. Using several such elbows a serpentine flow path can be created as in FIG. 3 (Elbows are also provided at the opposite ends of the tubes. These are not shown). Vertically oriented elbows may also be provided to join tubes on different racks 40.

It is therefore possible to create a single serpentine flow path through the entire set of tubes in a bioreactor. Of course, it may also be desirable to create two, three, or many more individual flow paths in a single bioreactor. Many different flow paths may be created, depending upon how the tubes are connected. It is also possible to use valves to create changeable flow paths. A pump is generally used to circulate the nutrient medium.

Since the microalgae growth depends on photosynthesis, carbon dioxide must be added to the circulating medium. It may also be desirable to remove the oxygen produced by the photosynthesis. FIG. 4 shows a simplified depiction of a device which can provide both of these functions. Gassing/degassing system 46 has housing 48. Two bioreactor tubes 42 are connected to housing 48. Inlet flow is provided through inlet 60. Outlet flow is provided through outlet 62. Thus, the interior of housing 48 is part of a flow path within the bioreactor.

Carbon dioxide inlet 50 introduces carbon dioxide. Oxygen outlet 52 allows the escape and collection of oxygen. It may also be desirable to maintain the circulating medium at a particular temperature. Thus, a heat exchange device is also provided. Aluminum helix 54 is a hollow tube. Coolant inlet 56 provides inlet cooling flow through the aluminum helix.

Coolant outlet carries away the coolant flow. The coolant used can be water which is cooled
160 by a separate chiller. Other coolants may of course be used as well.

Several gassing/degassing systems 46 can be installed at suitable locations within the
flow path of the bioreactor. Returning to FIG. 3, the reader will recall that simple elbows 44
may be used to direct the flow from one bioreactor tube 42 to another. Turning now to FIG.
4, those skilled in the art will realize that a gassing/degassing system 46 can be substituted for
165 any of the elbows (with suitable adjustment being made for the distance between inlet 60 and
outlet 62).

The bioreactor is largely a collection of simple components – such as a vertical rack
with multiple horizontal tubes in an appropriately spaced location. The connections between
many of the tubes will be made with elbows 44. The connection between other adjacent
170 tubes will be made using a gassing/degassing system 46. The “control and monitoring”
component is preferably part of gassing/degassing system 46. It is preferable to incorporate
numerous components in housing 48. For example, the housing can contain and/or mount:

- (1) carbon dioxide injecting systems;
- (2) oxygen removal systems;
- 175 (3) carbon dioxide sensors;
- (4) oxygen sensors;
- (5) pH sensors;
- (6) turbidity sensors;
- (7) flow sensors; and
- 180 (8) temperature sensors.

As explained previously, the housing may also contain a heat exchanger capable of
maintaining a desired temperature for the circulating medium. This would typically be a
liquid-to-liquid heat exchanger. However – in some ambient environments – it may be
possible to use a liquid-to-air exchanger. The systems for adding carbon dioxide and
185 removing oxygen are well known in the art and will thus not be described in detail. The same
may be said of the various sensors disclosed.

Space efficiency is a significant goal for the inventive photo-bioreactor. FIGs. 5A –
6B show a preferred embodiment that minimizes the amount of ground surface area required.
FIG. 5A shows how the components of photo-bioreactor 18 are generally supported by frame
190 38. The nutrient medium is collected in reservoir 68 near the top of the assembly. The

circulating medium flows out from the reservoir through reservoir outlet line 92 into inlet manifold 64. The inlet manifold feeds the liquid into multiple, independent flow paths.

In the embodiment shown, each independent flow path comprises a serpentine path of bioreactor tubes 42 connected by elbows 44. Each serpentine flow path creates one vertical "column" within the assembly. In this example there are fourteen such columns. Each column is fed circulating liquid by inlet manifold 64. Outlet manifold 66 collects the liquid as it exits each column. From the outlet manifold the liquid is fed to liquid pump 70. The liquid pump pressurizes the collected liquid medium and feeds it up through gasser tube 72 and back to reservoir 68.

Gas injector pump 76 takes in a desired gas through gas inlet 78. It pressurizes this gas beyond the pressure within the lower portion of gasser tube 72 and injects the gas into the liquid medium through gas injector 74. In the preferred embodiments the injected gas is simply ambient air. Thus, gas inlet 78 is configured to suck in ambient air – possibly using an appropriate filter to exclude dust and other particles.

The mixture of liquid medium and injected gas travels upward together within gasser tube 72. The gas and liquid remain in contact throughout this period. At the top of the gasser tube the contents empty into reservoir 68 and the circulation loop begins again.

FIG. 5B shows a perspective view of the same assembly. The reader should bear in mind that the depictions of the components are somewhat simplified. For example, the liquid and gas pumps are represented in a symbolic form. In addition, the bioreactor tubes and elbows are not necessarily drawn to the exact scale of a working unit.

FIG. 6A shows a front elevation view of the same assembly. In this view the user may easily perceive how the array of bioreactor tubes is arranged into vertical columns and horizontal rows. In this embodiment there are fourteen vertical columns 80 of bioreactor tubes. There are thirty-nine horizontal rows 82 of tubes in each vertical column (Other embodiments may include fifty or more rows in each vertical column). Each column represents an independent flow path. Each column is fed liquid medium through inlet manifold 64. The liquid medium then flows through the column's serpentine path until it reaches outlet manifold 66. In outlet manifold 66 the independent flow paths are reunited.

Pump intake line 88 takes the liquid from the outlet manifold to liquid pump 70. Of course, each column could have its own, separate return line to the liquid pump. The use of inlet manifold 64 and outlet manifold 66 represents only one way among many to create the

desired flow paths. In addition, the inlet and outlet manifolds would typically include valves allowing each column to be taken out of the circulation loop for harvesting of the biomass, cleaning, or some other purpose. These valves have not been shown for purposes of visual clarity. FIG. 6B provides a perspective view of the opposite end of the assembly (opposite to the end shown in FIG. 6A).

Returning to FIG. 5A, some additional features of the invention will be explained before turning to a discussion of dimensions for some of the preferred embodiments. A significant feature of the invention is its ability to promote the dissolving of gas into the circulating liquid by enhancing the gas/liquid interface. The injected gas is preferably air from the surrounding atmosphere. The percentage of carbon dioxide in this air will typically be around 0.04% by volume. The photo-bioreactor relies in part on carbon dioxide and it is therefore important to dissolve the available carbon dioxide into the water as efficiently as possible.

In prior art systems gaseous, concentrated carbon dioxide is injected into a liquid volume and allowed to "bubble through." The gas bubbles naturally rise and a free gas volume tends to accumulate at the top of any containment vessel used. This fact means that a significant volume of gas is surrounded only by other gas and has no opportunity to contact the liquid. The present invention reduces this problem and increases the rate at which the gas dissolves into the liquid.

Looking again at FIG. 5A, those skilled in the art will realize that the maximum pressure within the circulating system occurs at pump discharge line 90. The gas is injected very near this point of maximum pressure (Note location of gas injector 74). Further, the bubble size of the injected gas is controlled by using a suitable mesh or array of orifices. FIG. 7 shows the use of an exemplary mesh 86 across the injection opening. With appropriate injection pressure, this mesh produces small bubbles in the range of 0.5 mm to 1.0 mm. A smaller bubble creates a larger ratio of bubble surface area to bubble volume. This larger ratio enhances the solution rate.

Returning now to FIG. 5A, the reader will also observe how the gas bubbles are introduced at the bottom of a tall vertical column (gasser tube 72). This column may have a height of 15 meters or more. The gas bubbles flow upward with the ascending liquid. This maximizes the contact time between the gas and the liquid and prevents the formation of a large gas volume (though some aggregation of the bubbles will occur). Those skilled in the

255 art will know that the bubble ascent rate is a strong function of bubble size, and that smaller bubbles ascend more slowly. The use of an injector that limits bubble size therefore increases the amount of time that the gas bubbles remain in gasser tube 72 (in addition to improving the surface area to volume ratio for each bubble).

The flow rate of the mixture within the gasser tube is a function of the volumetric
260 flow of liquid pump 70 and the cross-sectional area of the gasser tube. A larger gasser tube produces a slower flow rate. Once the contents of the gasser tube reach the top they flow into reservoir 68. At this point any gas remaining will tend to migrate to the surface. Thus, it is preferable to design the components so that substantially all the injected gas is dissolved before the mixture reaches the reservoir. This may be done by matching the liquid flow rate
265 in the gasser tube fairly closely to the bubble ascent rate, although some difference in the two velocities is desirable to promote turbulent flow.

Of course, in order to maximize the dissolution of the gas in the liquid, an excess amount of gas is preferable. Excess gas presents no problem for the preferred embodiments of the invention since they are injecting atmospheric air. Any excess gas may be vented back
270 to the surrounding atmosphere.

The reader's understanding may benefit from some detailed explanation of the dimensions used in some of the preferred embodiments. In creating such a system, one may consider the following things, among others:

1. The diameter of the bioreactor tubes;
- 275 2. The overall length of each flow path;
3. The spacing of the bioreactor tubes (in order to ensure adequate sunlight exposure);
4. The flow rate within each flow path;
5. The dimensions of the gasser tube;
- 280 6. The flow rate within the gasser tube;
7. The injection of the gas; and
8. The overall "footprint" of the complete assembly.

A pilot scale photo-bioreactor has been developed and its dimensions are provided. The reader should bear in mind that both smaller and larger embodiments are possible. The
285 pilot scale embodiment uses 3710 total meters of transparent PVC tubes. The tubes are given UV radiation protection. As shown in FIGs. 5A – 6B, the tubes are arranged into vertical

columns and horizontal rows. The tubes in each column are joined together by the elbows to create a serpentine flow path lying in a vertical plane. In a preferred embodiment, each independent flow path is 265 meters long. The external support frame 38 includes numerous cross pieces that support the tubes along their length (Most of these are not shown for purposes of visual clarity).

A preferred embodiment includes 14 vertical columns with 53 tubes in each column. Each of these tubes has an internal diameter of 5 cm. The straight vertical portion of gasser tube 72 is 8 - 16 meters long. It has an internal diameter of 10 cm. Each of the transparent tubes has a cross-sectional area of 19.6 square centimeters. Since there are 14 separate flow paths in this exemplary photo-bioreactor the combined cross-sectional area for all the tubes is 275 square centimeters. The cross-sectional area in the gasser tube is 78.5 square centimeters. Thus, the flow in each transparent tube is considerably slower than the flow in the gasser tube.

The entire assembly of FIGs. 5 - 6 occupies a compact volume. It is approximately 5 m long by 2 m wide by 10 m high. Despite its compact size, the inventive photo-bioreactor is capable of cultivating approximately 10,000 L of microalgae medium while using only 10 square meters of total surface area.

The components are depicted in somewhat-simplified form. The pump, for example, is not shown in detail. This component is preferably a diaphragm unit rather than a centrifugal one, as centrifugal pumps tends to harm the growing microalgae. In addition, numerous conventional components have not been depicted. For instance, it is desirable to include drain valves that allows the medium to be removed from the photo-bioreactor for filtering, drying, and further processing. Such valves may be located in the vicinity of outlet manifold 66, but they have not been illustrated. As explained in the following, it may also be necessary to include gas collection and removal chambers.

The inventive assembly maximizes the contact area and contact duration between the injected gas and the liquid medium. This feature increases the solution rate of the gas into the liquid medium and represents a significant advantage. A comparison of the inventive system to the prior art will serve to illustrate this advantage.

The prior art approach is to inject carbon dioxide into horizontal, tubular manifolds or large liquid holding tanks. In the case of a horizontal manifold, the injected gas tends to

aggregate quickly into a layer at the top of the manifold. In the case of a holding tank the gas tends to bubble quickly through the liquid and then aggregate at the top or escape altogether.

320 A typical holding tank is about 1.5 meters deep. See, for example, the tanks 121-129 at Figure 12 of U.S. Publication No. 2010/0159579. If gas bubbles are injected conventionally they tend to create an initial bubble size of about 0.75 cm or more. The bubble ascent rate for the smaller bubbles in this range is approximated by the expression:

$$v = \frac{1}{9} \left(\frac{g}{\nu} \right)^2 d^2$$

where d is the bubble diameter, g is gravitational acceleration and ν is

325 the kinematic viscosity of water. Using this expression, a gas bubble having a diameter of 0.75 cm will ascend at a rate of about 1.1 m/s. Such a bubble will pass through a 1.5 meter deep tank in only 1.4 seconds.

The vertically-oriented gasser tube in the present invention produces a very different result. Liquid pump 70 is configured to produce a flow rate of about 3.5 cubic meters per hour. The gasser tube has an internal diameter of 10 cm in this example, producing a cross
330 sectional area of 0.00785 square meters. Pumping 3.5 cubic meters per hour into this tube produces a modest linear flow rate of only 0.124 m/s.

An exemplary embodiment uses a gasser tube that is 16 meters high. This fact means that it takes 129 seconds (over two minutes) for the liquid flowing at 0.124 m/s to flow from
335 the bottom of the gasser tube to the top.

Of course, gas is also being injected into the gasser tube. Returning to FIG. 5, the reader will recall that gas injector pump 76 injects a gas near the bottom of the gasser tube. In this example, the gas is air. The gas is injected at the rate of 2 cubic feet per minute (0.057 cubic meters per minute or about 3.5 cubic meters per hour). Thus, the liquid and gas
340 injection rates into the bottom of the gasser tube are about equal. The total flow in the injector tube is around 7 cubic meters per hour (gas and liquid combined) so the actual average velocity within the gasser tube is about 0.25 m/s. Even so, the "dwell" of the liquid/gas mixture within the 16 m high tube is still $16/0.25 = 64$ seconds.

Further, the 1-minute-plus dwell is further enhanced by a substantially reduced gas
345 bubble size. The injection mesh shown in FIG. 7 limits the bubble size of the injected gas to between 0.5 mm and 1.0 mm. A differential pressure between the injected gas and the liquid in the gasser tube of 1 to 4 bar is used. Assuming an average bubble size of 0.75 mm, the bubble ascent rate can be approximated as:

$$v = \frac{1}{3} \left(\frac{d}{2} \right)^2 g / \nu$$

350 Using this expression, a gas bubble having a diameter of 0.75 mm will ascend at the rate of only 0.033 m/s. This ascent rate will be added to the ascent rate of the mixture as a whole. Very roughly speaking, the gas bubbles within the mixture within the gasser tube will ascend at about 0.28 m/s, whereas the liquid will ascend at about 0.22 m/s. This is enough velocity difference to create turbulent mixing and enhance the solution of the gas into the
355 liquid.

The small bubble size produces a much larger surface-area-to-mass ratio for the gas contained in each bubble. Any volume of gas within the gasser tube is very near a liquid-to-gas interface surface at all times. The gas is subjected to this state for about one full minute. By the time the gas has reached the top of the gasser tube, the liquid is preferably saturated.
360 Some excess gas may remain and this simply bubbles out of the reservoir. The reader will recall that – in this example – the gas is simply ambient air. The escape of some of this air is therefore not a problem.

Prior art systems create a poor scenario for dissolving the gas into the liquid. Most of the gas winds up being segregated into a large gas volume. Further, the “dwell” time of the
365 gas bubbles in the liquid may be only 1 to 2 seconds. For these reasons, the prior art systems have been forced to use concentrated carbon dioxide as the feed gas for a photo-bioreactor.

In contrast, the approach taken in the present invention prevents the formation of large segregated gas volumes, maximizes the surface-area-to-volume ratio for the bubbles, and provides a “bubble-through” time of 1 minute or even more. With these advantages, the
370 present invention is able to use ambient air as the feed gas. To be sure, the present invention could also use concentrated carbon dioxide. However, its ability to feed the photo-bioreactor using air taken from the surrounding atmosphere is significant.

Some additional details of the various possible embodiments are listed in the following:

- 375 1. The reservoir need not be very large. It can be practically any size and shape. There are operational advantages to mounting it on the top of the photo-bioreactor but it may be located in other places;
2. The gasser tube does not have to be round and it does not have to have a constant cross-section;

380 3. The bubble-size limiter used in the air injection may be a plate with many holes rather than a wire mesh;

4. The photo-bioreactor can be located near a source of carbon-dioxide pollution, such as a coal-fired power plant. However, it can be located anywhere there is available space and sunlight;

385 5. It is preferable to match the ascent rate of the liquid nutrient medium and the gas bubbles in the gasser tube so that they are approximately equal, meaning that they are within 0.12 meters per second and even more preferably within .08 m/s; and

6. It is preferable to match the volume of liquid nutrient medium and injected gas within the gasser tube so that the volume of one is within 50% of the volume of the other.

EXAMPLE OF OPERATION

The photo-bioreactor illustrated and described may be operated in a variety of ways to accomplish differing results. The following descriptions should not be viewed as limiting in any way. The use of the photo-bioreactor for the production of biodiesel has been described
395 previously. This process generally involves the promotion of green algae growth to a stable level, followed by filtering and drying. The dried biomass may then be pressed and processed to produce biodiesel and other products in ways known to those skilled in the art.

The photo-bioreactor may also be used for the production of hydrogen. Hydrogen is a natural – albeit transient – product of several microbial driven biochemical reactions. The
400 hydrogen is produced mainly in anaerobic fermentation processes. In addition, certain microorganisms produce enzymes that can catalyze hydrogen synthesis if an outside energy source, like sunlight, is available. The known bio-hydrogen production processes are: (1) Biophotolysis of water using green algae and blue algae (cyanobacteria), through a direct or indirect process; (2) Photofermentation; (3) Dark fermentation; and (4) Hybrid systems
405 combining one or more of these processes.

In this context microalgae could provide several types of different biofuels, including: (1) Microalgae-derived biodiesel; (2) Methane produced through anaerobic digestion of microalgae biomass after lipid extraction; (3) Hydrogen produced by water photolysis during photosynthesis; and (4) Ethanol produced from microalgae biomass after lipid extraction
410 (which is still expected to contain a large carbohydrate mass fraction for fermentation).

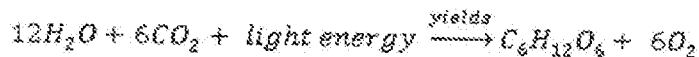
Direct biophotolysis is the dissociation of the water molecule due to the action of light energy. This process occurs naturally during green algae photosynthesis. However, the concurrent production of oxygen strongly inhibits the enzyme hydrogenase that catalyzes the production of hydrogen. Therefore, anaerobic conditions are essential for hydrogen
415 production in larger quantities. For large-scale production, the so-called indirect biophotolysis processes have been proposed, in which carbon dioxide is first fixed into carbohydrates and then used in a separate step to produce hydrogen.

Both direct biophotolysis and indirect biophotolysis with filamentous heterocystous cyanobacteria show simultaneous production of oxygen and hydrogen. Either approach
420 would require expensive gas separation. On the other hand, reversible hydrogenase-based indirect biophotolysis processes have, at least conceptually, major advantages over the nitrogenase-based systems. The specific hydrogen evolution activities of reversible hydrogenases are almost a thousand-fold higher than those of nitrogenase and, most importantly, require no expensive adenosine tri-phosphates. Thus, this approach appears to
425 have real practical potential.

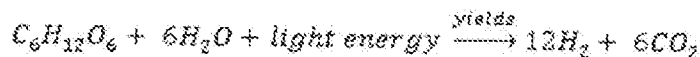
Hydrogen production by biophotolysis could be defined as the dissociation of the water molecule through the action of light energy. In the process, sulfur is a key component of the amino acids for the proteins where oxygen is produced during the photosynthesis. Therefore, if one wants to inhibit oxygen production then sulfur nutrient-deprived green algae
430 cultures have good potential for hydrogen production in anaerobic conditions. This is true since the enzyme Fe-hydrogenase, which is responsible for the hydrogen production process using two electrons brought by the protein ferredoxin ($2H^+ + 2Fd^- \xrightarrow{\text{yields}} H_2 + 2Fd$) has its activity inhibited by oxygen. However, photosynthesis should not be inhibited in the first stage of the process, so that the desired microalgae can grow and produce biomass.

Hence, the indirect biophotolysis process in a bioreactor should be divided into two
435 separate stages. In the first stage, aerobic conditions (air injection) are applied to increase the biomass up to stabilization. In the second stage, the air supply is cut off and the process continues under anaerobic conditions. In order to use a photo-bioreactor such as shown at FIGs. 5 and 6 for this two-stage process, a cycling regime should be established -- as will be
440 explained.

The following reaction summarizes the aerobic first stage of the indirect biophotolysis process:



The following reaction summarizes the anaerobic second stage of the indirect
445 biophotolysis process:



Under this approach, the photo-bioreactor such as shown in FIG. 5 will not be operated in a steady state. Rather, it will first be operated in an aerobic stage to grow the biomass in the presence of air-injection and with oxygen as a significant, circulating product.
450 In the second stage the air supply will be cut off and free hydrogen will be produced.

Even within the stages the reaction rates will vary according to sunlight and temperature. Sunlight only falls on the photo-bioreactor during daylight hours. Even during daylight hours the reaction rates vary with ambient temperature.

The aerobic stage of the process typically runs for about 8 days, at which point the
455 increase rate for the biomass has tapered off. The oxygen concentration is also stable at this point. When the system is switched to the anaerobic stage, the air supply is cut off and the algae perform only mitochondrial respiration. From this point forward the biomass is consumed and its mass fraction decreases. Likewise, the mass fraction of the oxygen decreases during the anaerobic stage since it is consumed by mitochondrial respiration.

460 In the aerobic stage the carbon dioxide mass fraction remains fairly constant and fairly low (since the carbon dioxide being injected is consumed by the photosynthesis process). In the anaerobic stage the air supply is cut off and the carbon dioxide mass fraction increases.

Hydrogen production is inhibited during the aerobic stage by the absorbed oxygen in
465 the medium, which inhibits the hydrogenase enzyme activity. In fact, the hydrogen mass fraction is practically zero.

However, in the anaerobic stage, as the oxygen starts to be consumed by the microalgae mitochondrial respiration the hydrogen mass fraction will rise. The hydrogenase enzyme starts to catalyze the reaction shown previously. However, hydrogen will only be
470 produced in the presence of carbohydrates ($C_6H_{12}O_6$). Essentially, the carbon dioxide in the injected air is first "fixed" into carbohydrates and then used in the second stage to produce

hydrogen. The hydrogen produced bubbles out of the circulating liquid as a gas. It may then be collected, separated, and stored.

The reader will thus appreciate that the present invention provides a comprehensive
475 and space-efficient system for producing biodiesel, hydrogen, or potentially other bio-fuels
from microalgae. The foregoing description and drawings comprise illustrative embodiments
of the present invention. Having thus described exemplary embodiments of the present
invention, it should be noted by those skilled in the art that the within disclosures are
exemplary only, and that various other alternatives, adaptations, and modifications may be
480 made within the scope of the present invention.

Many modifications and other embodiments of the invention will come to mind to one
skilled in the art to which this invention pertains having the benefit of the teachings presented
in the foregoing descriptions and the associated drawings.

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CLAIMS

505 Having described my invention, I claim:

Claim 1. A photo-bioreactor for growing biomass within a liquid nutrient medium, comprising:

- a. a support frame, having a top and a bottom;
- b. a plurality of separate flow paths supported by said support frame, each
510 of said separate flow paths including a plurality of connected transparent tubes;
- c. a reservoir connected to said plurality of separate flow paths, said reservoir configured to feed said liquid nutrient medium to said separate flow paths, said reservoir being located proximate said top of
515 said support frame;
- d. a liquid pump, having an intake side and a discharge side, said liquid pump being located proximate said bottom of said support frame;
- e. said intake side of said liquid pump being configured to take in said liquid nutrient medium from said separate flow paths;
- f. a gasser tube, said gasser tube being vertically oriented and having a
520 lower portion and an upper portion;
- g. said lower portion of said gasser tube being connected to said discharge side of said liquid pump;
- h. said upper portion of said gasser tube being connected to said
525 reservoir;
- i. a gas injector configured to inject a gas into said lower portion of said gasser tube, thereby creating a mixture of said liquid nutrient medium and said gas in said gasser tube;
- j. said gas injector including a bubble size limiter configured to limit an
530 average size of a gas bubble introduced by said gas injector into said liquid nutrient medium to below 2 millimeters; and
- k. said gasser tube, said liquid pump, and said gas injector being configured so that said liquid nutrient medium requires at least 30 seconds to travel from said bottom of said gasser tube to said top of
535 said gasser tube.

- Claim 2. A photo-bioreactor for growing biomass within a liquid nutrient medium as recited in claim 1, wherein said gas is air.
- 540 Claim 3. A photo-bioreactor for growing biomass within a liquid nutrient medium as recited in claim 2, further comprising:
- a. an air pump configured to feed pressurized air to said air injector; and
 - b. wherein said air pump pulls in ambient air.
- 545 Claim 4. A photo-bioreactor for growing biomass within a liquid nutrient medium as recited in claim 2, further comprising an air pump configured to feed pressurized air to said air injector.
- Claim 5. A photo-bioreactor for growing biomass within a liquid nutrient medium as
550 recited in claim 1, wherein said bubble size limiter is a wire mesh.
- Claim 6. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 1, wherein said gasser tube has a height greater than 10
555 meters.
- Claim 7. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 1, wherein said liquid pump and said gas injector are
configured such that an ascent rate of said liquid nutrient medium and an
ascent rate of said gas bubbles in said gasser tube are approximately equal.
560
- Claim 8. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 7, wherein said liquid pump and said gas injector are
configured such that an ascent rate of said liquid nutrient medium and an
ascent rate of said gas bubbles in said gasser tube are within 0.08 meters per
565 second of each other.

Claim 9. A photo-bioreactor for growing biomass within a liquid nutrient medium as recited in claim 1, wherein a volume of liquid nutrient medium pumped into said gasser tube by said liquid pump and a volume of said gas injected into said gasser tube by said gas injector are approximately equal.

Claim 10. A photo-bioreactor for growing biomass within a liquid nutrient medium as recited in claim 9, wherein said liquid pump pumps said liquid nutrient medium at a rate between about 3 cubic meters per hour and about four cubic meters per hour.

Claim 11. A photo-bioreactor for growing biomass within a liquid nutrient medium, comprising:

- a. a support frame, having a top and a bottom;
- b. a plurality of hollow, transparent tubes defining a flow path through which said liquid nutrient medium circulates, said tubes being supported by said support frame;
- c. a liquid pump, having an intake side and a discharge side, said liquid pump being located proximate said bottom of said support frame;
- d. said intake side of said liquid pump being configured to take in said liquid nutrient medium from said plurality of transparent tubes;
- e. a gasser tube, said gasser tube being vertically oriented and having a lower portion and an upper portion;
- f. said lower portion of said gasser tube being connected to said discharge side of said liquid pump;
- g. said upper portion of said gasser tube being connected to said plurality of transparent tubes;
- h. a gas injector configured to inject a gas into said lower portion of said gasser tube, thereby creating a mixture of said liquid nutrient medium and said gas in said gasser tube;
- i. said gas injector including a bubble size limiter; and
- j. said gasser tube, said liquid pump, and said gas injector being configured so that said liquid nutrient medium requires at least 30

seconds to travel from said bottom of said gasser tube to said top of
600 said gasser tube.

Claim 12. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 11, wherein said gas is air.

605 Claim 13. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 12, further comprising:
a. an air pump configured to feed pressurized air to said air injector; and
b. wherein said air pump pulls in ambient air.

610 Claim 14. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 12, further comprising an air pump configured to feed
pressurized air to said air injector.

Claim 15. A photo-bioreactor for growing biomass within a liquid nutrient medium as
615 recited in claim 11, wherein said bubble size limiter is a wire mesh.

Claim 16. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 11, wherein said gasser tube has a height greater than 10
meters.

620 Claim 17. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 11, wherein said liquid pump and said gas injector are
configured such that an ascent rate of said liquid nutrient medium and an
ascent rate of said gas bubbles in said gasser tube are approximately equal.

625 Claim 18. A photo-bioreactor for growing biomass within a liquid nutrient medium as
recited in claim 17, wherein said liquid pump and said gas injector are
configured such that an ascent rate of said liquid nutrient medium and an
ascent rate of said gas bubbles in said gasser tube are within 0.08 meters per
630 second of each other.

Claim 19. A photo-bioreactor for growing biomass within a liquid nutrient medium as recited in claim 11, wherein a volume of liquid nutrient medium pumped into said gasser tube by said liquid pump and a volume of said gas injected into said gasser tube by said gas injector are approximately equal.

Claim 20. A photo-bioreactor for growing biomass within a liquid nutrient medium as recited in claim 19, wherein said liquid pump pumps said liquid nutrient medium at a rate between about 3 cubic meters per hour and about four cubic meters per hour.

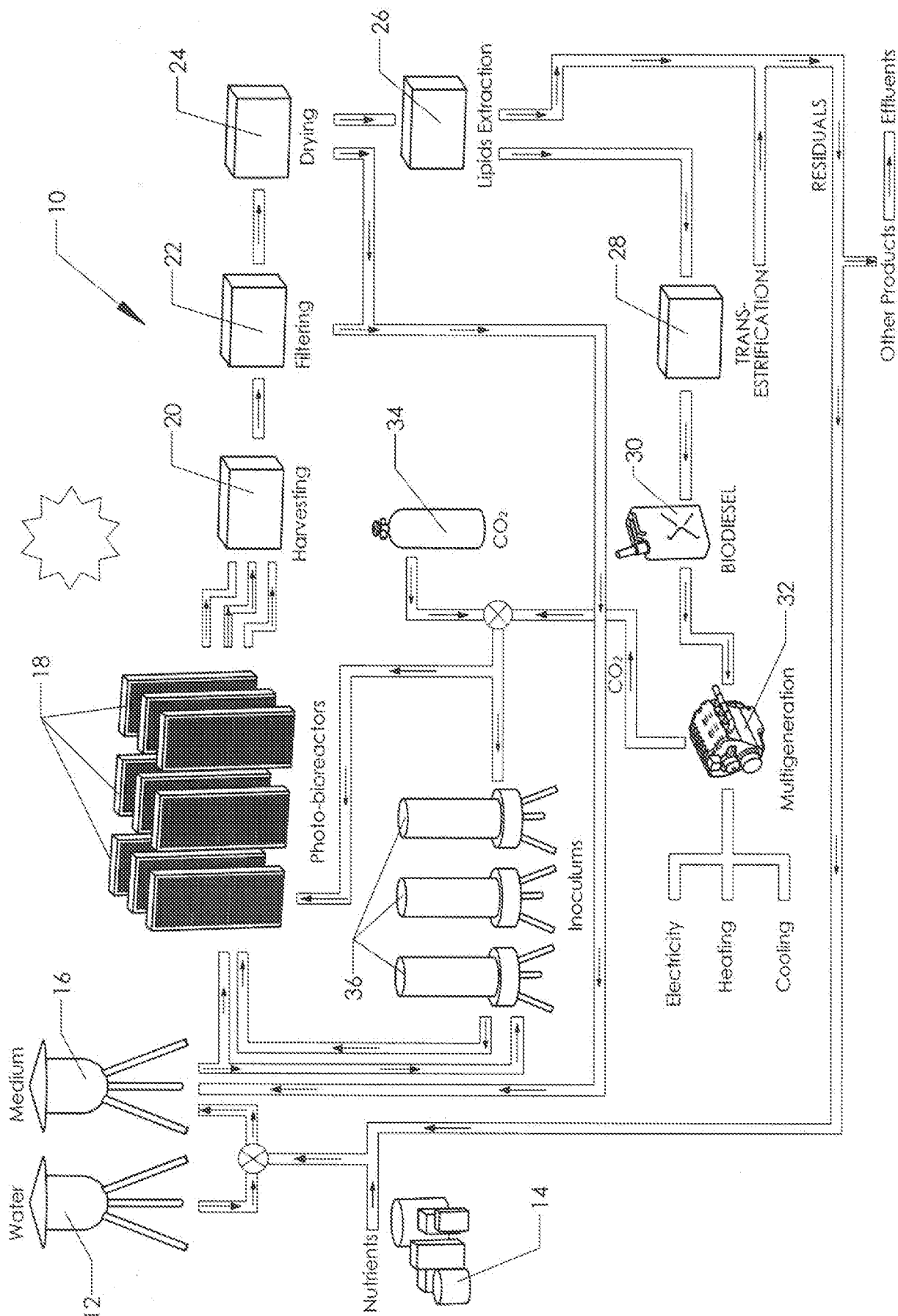


FIG. 1

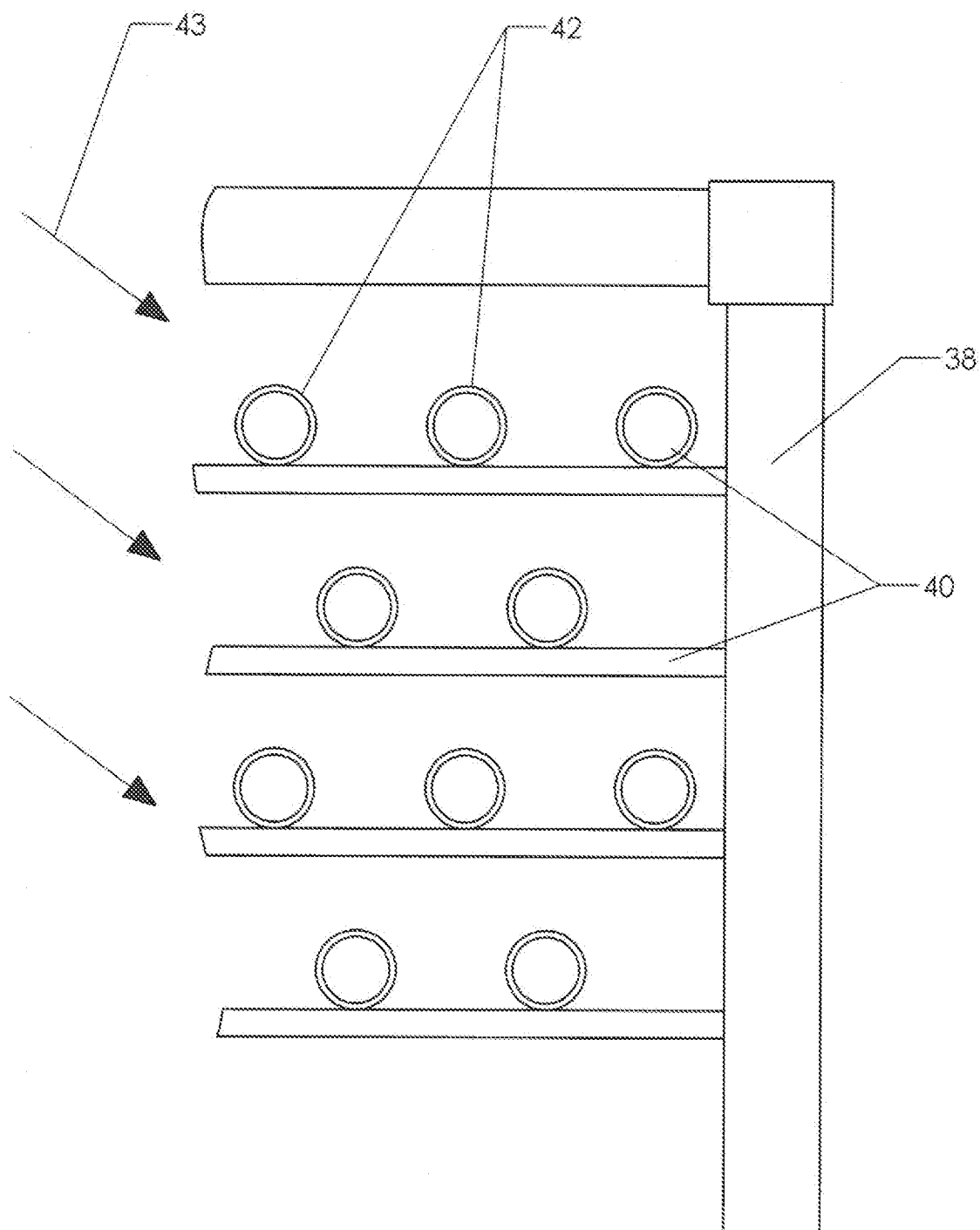


FIG. 2

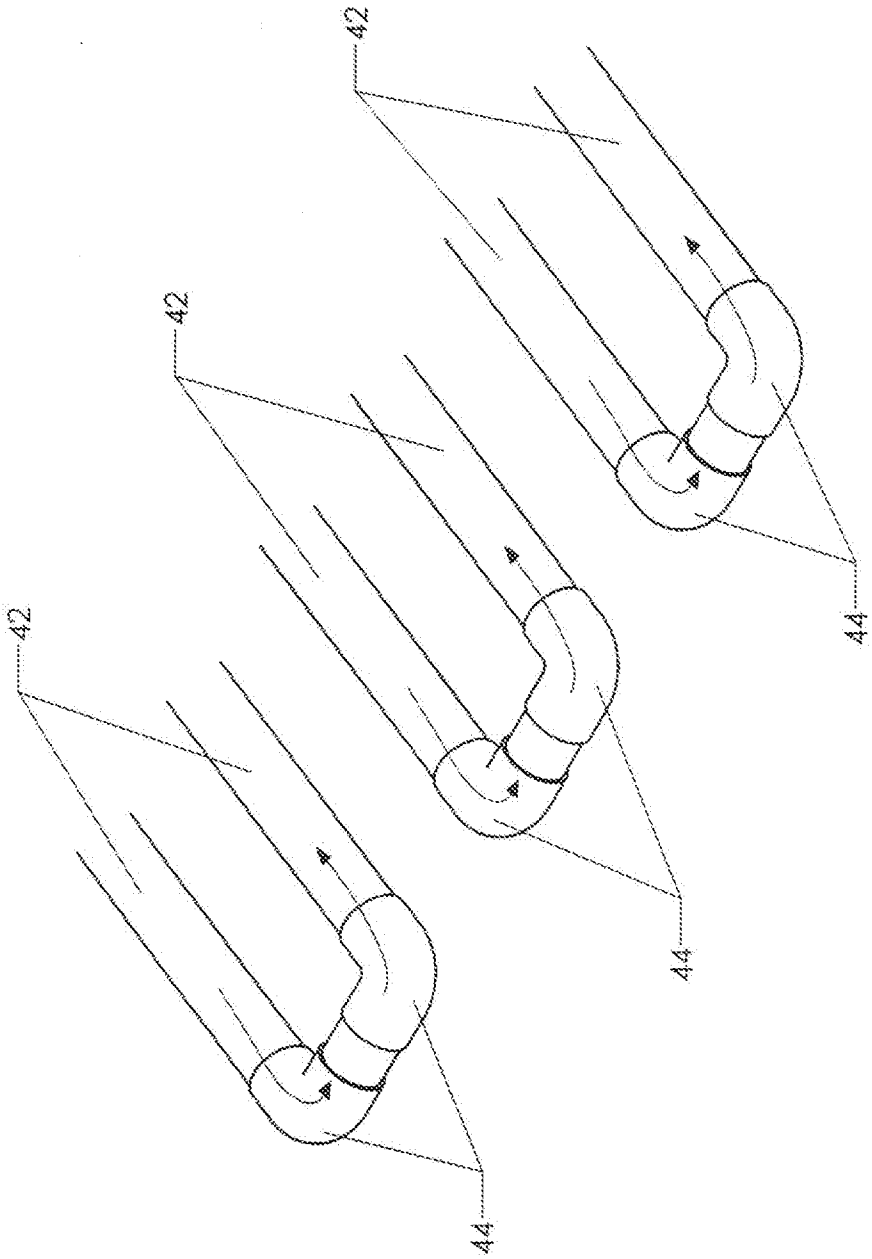


FIG. 3

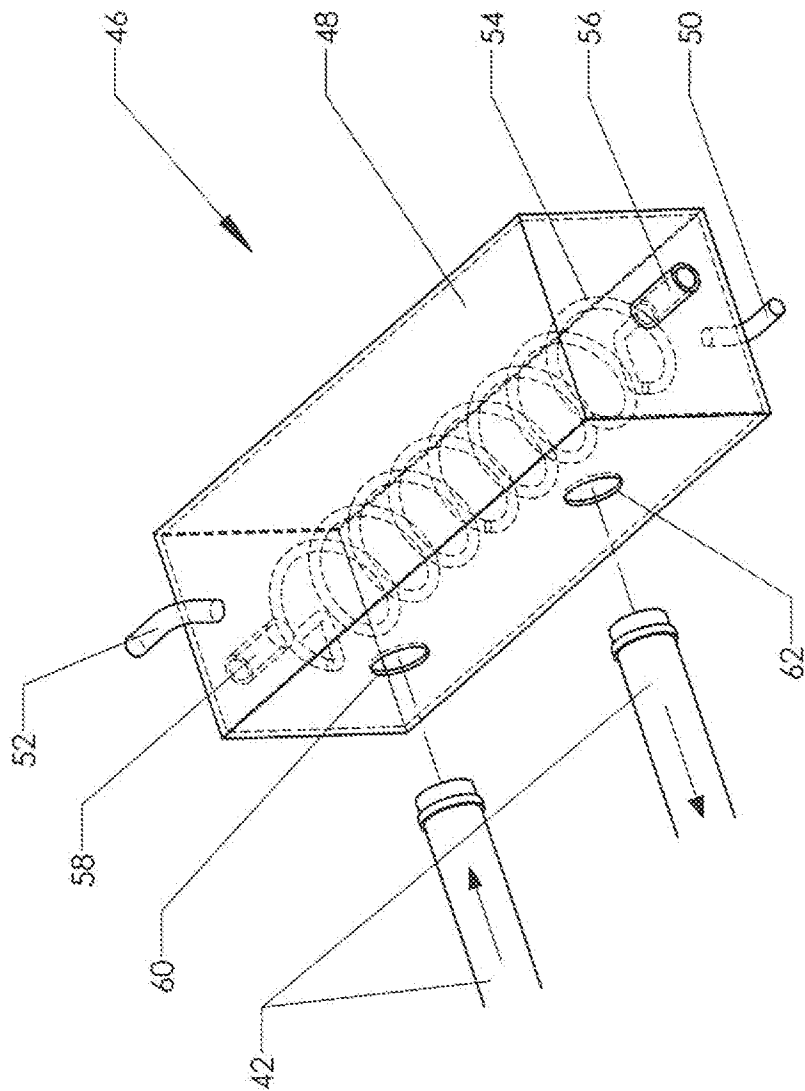


FIG. 4

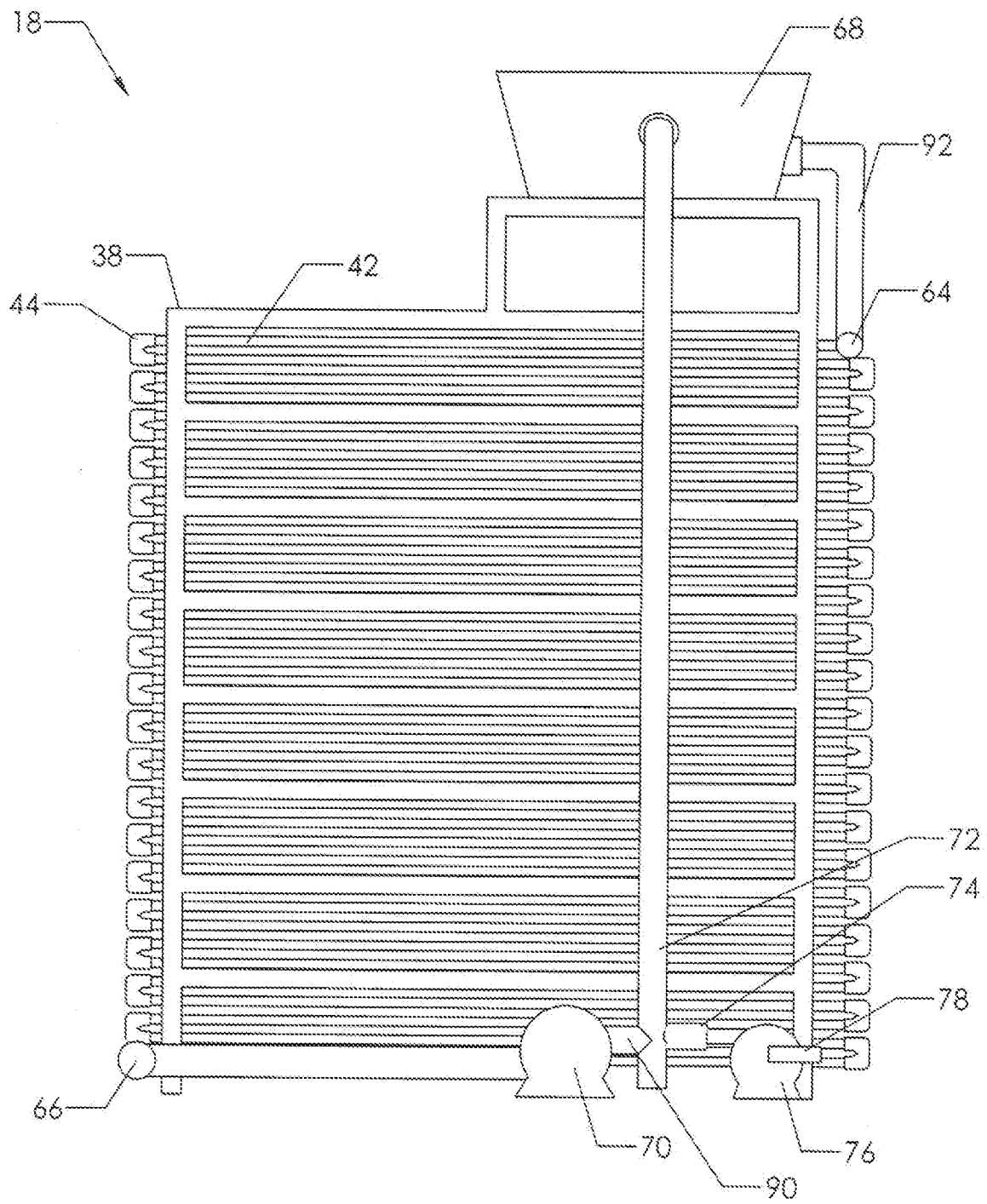


FIG. 5A

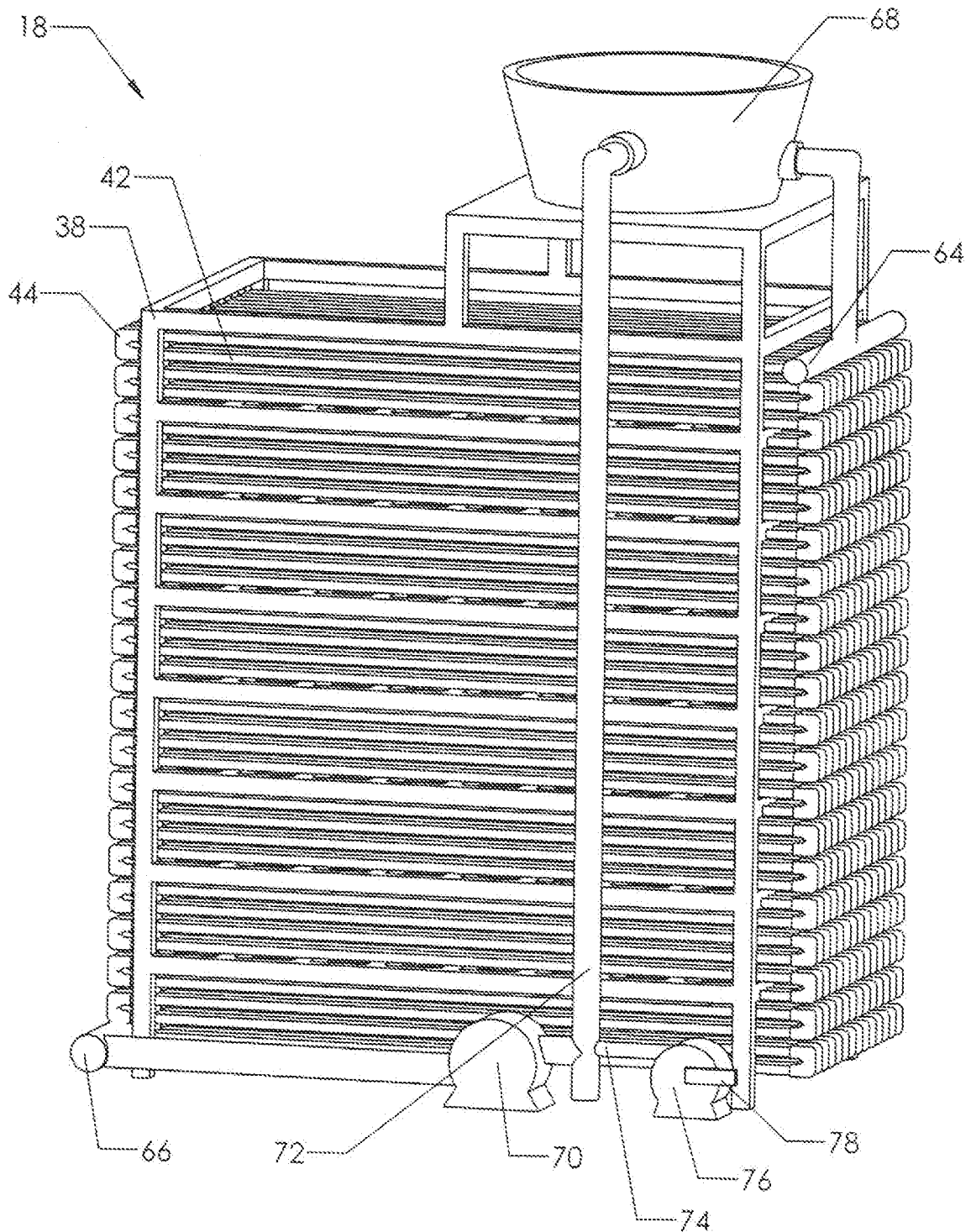


FIG. 5B

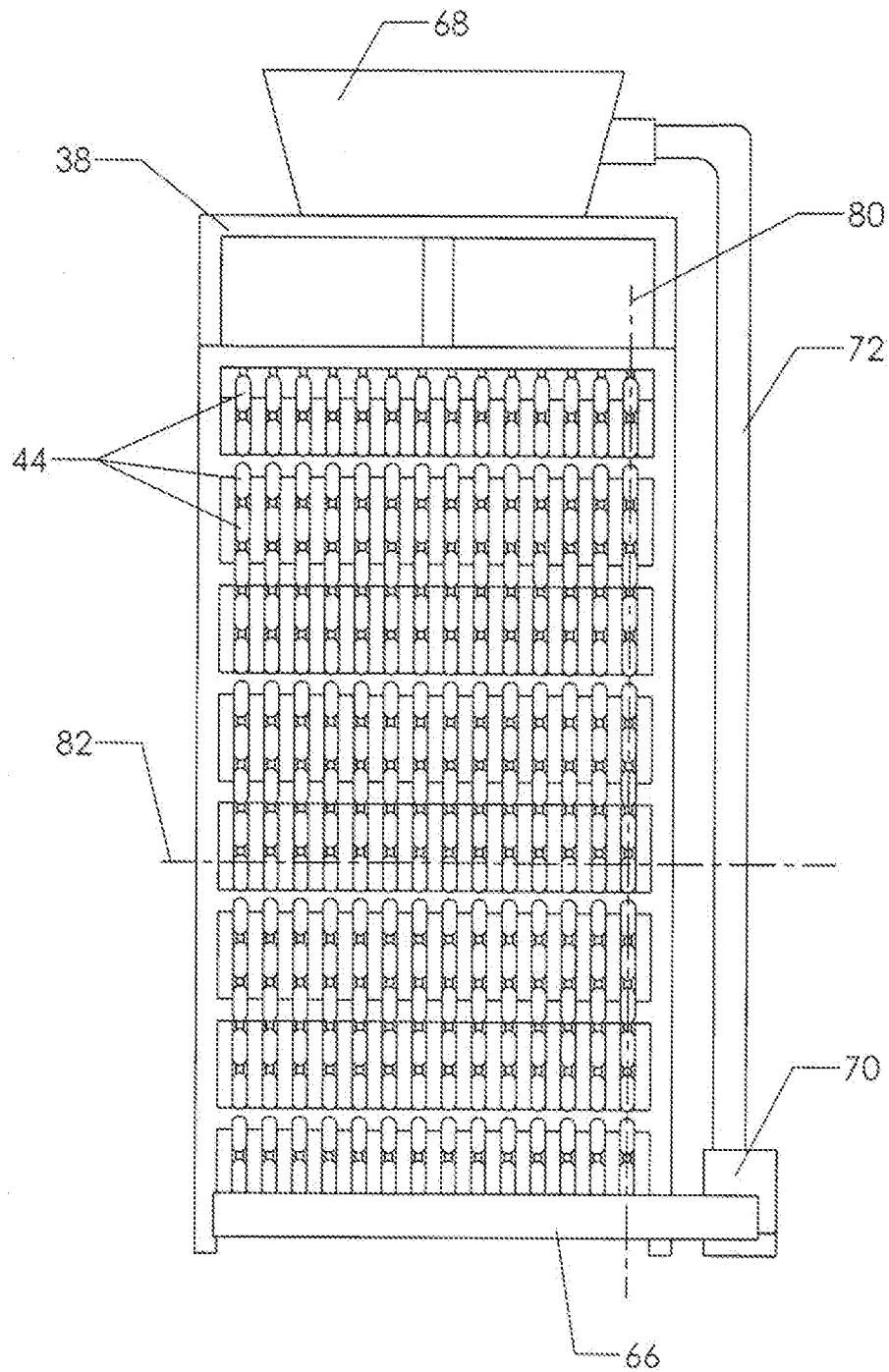


FIG. 6A

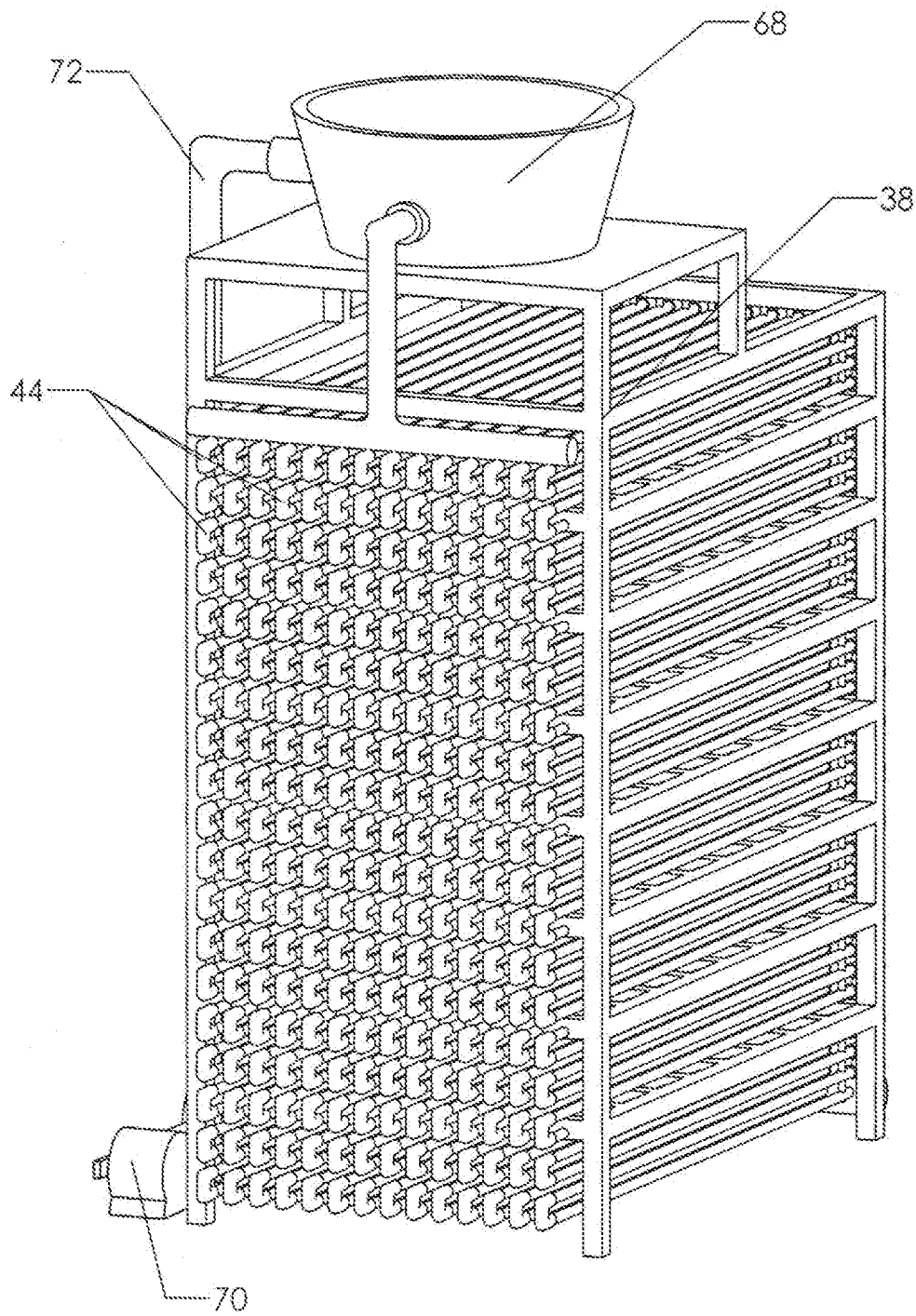


FIG. 6B

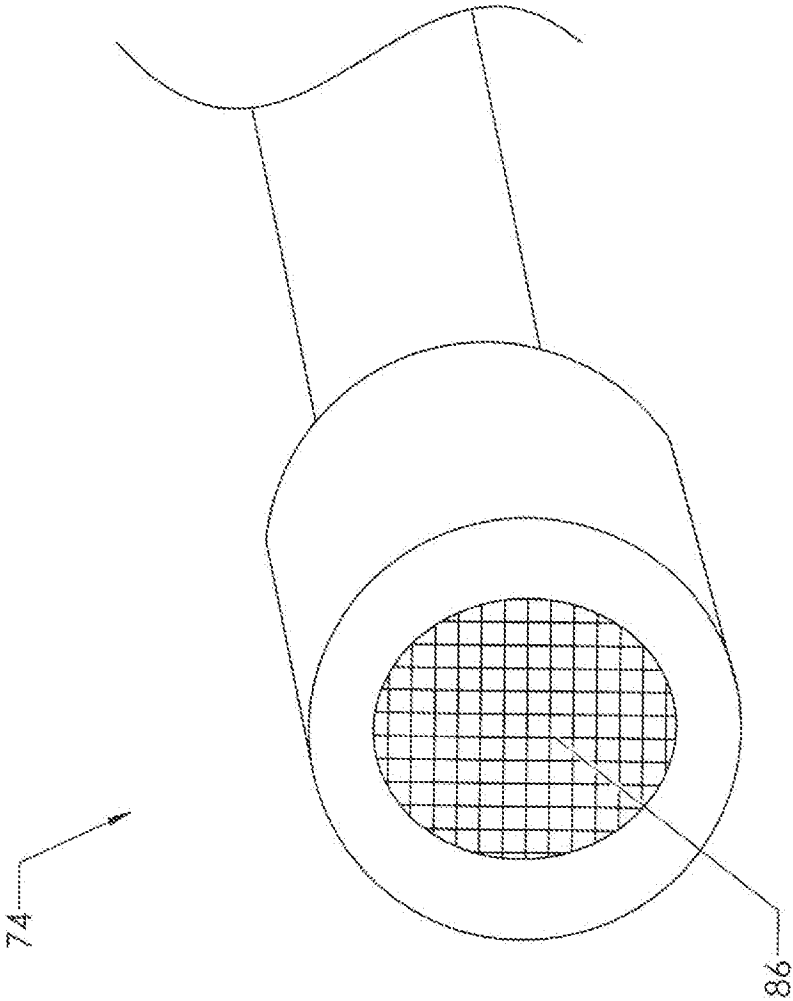


FIG. 7

INTERNATIONAL SEARCH REPORT		International application No. PCT/US16/37154		
A. CLASSIFICATION OF SUBJECT MATTER IPC: C12M 1/00(2006.01),1/26(2006.01) USPC: 435/292.1 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) U.S. : 435/292.1 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) Please See Continuation Sheet				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.		
Y	US 2010/0159579 A1 (SCHURING et al.) 24 June 2010 (24.06.2010), entire document.	1-20		
Y	US 2008/0032396 A1 (CHOKSHI) 07 February 2008 (07.02.2008), entire document.	1-20		
Y	Emery et al. Oxygenation of Intensive Cell-Culture System, Appl Microbiol Biotechnol (1995), 43: 1028-1033.	5 and 15		
☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.				
<table style="width: 100%; border: none;"> <tr> <td style="width: 50%; vertical-align: top;"> * Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed </td> <td style="width: 50%; vertical-align: top;"> "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family </td> </tr> </table>			* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family			
Date of the actual completion of the international search 05 July 2016 (05.07.2016)		Date of mailing of the international search report 12 JUL 2016		
Name and mailing address of the ISA/US Mail Stop PCT, Attn: ISA/US Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450 Facsimile No. (571) 273-3201		Authorized officer John Nguyen Telephone No. (571) 272-4300		

INTERNATIONAL SEARCH REPORT	International application No. PCT/US16/37154
Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)	
This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:	
1.	☐ Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
2.	☐ Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3.	☐ Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).
Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)	
This International Searching Authority found multiple inventions in this international application, as follows:	
1.	☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2.	☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of any additional fees.
3.	☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4.	☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
Remark on Protest	☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee. ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation. ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US16/37154

Continuation of B. FIELDS SEARCHED Item 3:

EAST: US-PGPUB; USPAT; FPRS; EPO; JPO; DERWENT

Search Terms: reactor, bioreactor, photobioreactor, tubes, support, frame, nutrient, oxygen, carbon, dioxide, heat, exchang\$3, bubble, size, filter, mesh, limit, gas, air, inject\$3, diffus\$4, injector, wire