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(54) Title: WATER-FREE SURFACE SIZING COMPOSITION AND METHOD FOR TREATING A PAPER SUBSTRATE WITH SAME

(57) Abstract: Water-free compositions suitable for application as a surface size to a cellulosic substrate, and methods of applying the water-free compositions to the surface of a cellulosic substrate.



WATER-FREE SURFACE SIZING COMPOSITION AND METHOD FOR TREATING A PAPER SUBSTRATE WITH SAME

[0001] This application claims the benefit of U.S. Provisional Patent Application No. 61/834,530, filed June 13, 2013, the disclosure of which is herein incorporated by reference in its entirety.

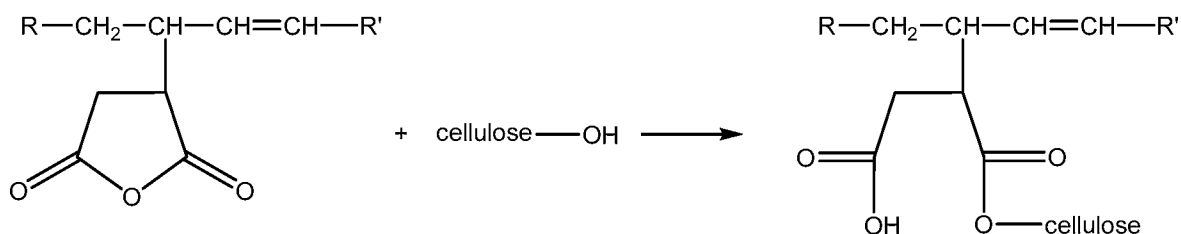
FIELD

[0002] The technical field relates to compositions containing sizing agents useful in the paper industry and to a method of treating a paper substrate with such compositions.

BACKGROUND

[0003] Cellulose, which is the main component of paper substrates, is hydrophilic and polar. These characteristics result in rapid water penetration in the paper substrate. This phenomenon can be delayed by hydrophobation of the paper substrate. This operation is known as “paper sizing.” By this operation, the penetration of polar liquids (*e.g.*, water or inks) in the paper is delayed by recovering the cellulose fibers with a hydrophobic substance, which is called a sizing agent. The sizing agent is usually added to the paper pulp and retained by the cellulose fibers in the wet end part of the paper manufacturing process. This is called internal sizing. However, the sizing agent can also be applied to the surface of the dried or partially dried paper, *i.e.*, surface sizing.

[0004] Typical sizing agents include rosin, alkenyl succinic anhydrides (“ASAs”) and alkyl ketene dimers (“AKDs”). ASAs are good candidates for surface sizing of paper substrates due to their high reactivity towards the hydroxyl groups of the cellulose. The reaction between ASAs and cellulose can be represented as follows:



The formation of covalent bonds between the cellulose and the ASA translates into an efficient sizing and the resulting paper products show a good resistance to polar liquids penetration.

[0005] For surface sizing applications, the ASA is generally emulsified in water and the emulsion is applied using a sizing press or a coater. ASA emulsion can also, but less often, be applied using a shower. However, some curl problems have been observed when applying ASA water emulsions using showers. Moreover, for efficiency reasons and to obtain a sizing as homogeneous as possible, the particle size has to be controlled and a limited particle size range must be obtained. The water emulsion containing ASA must also be used quickly so as to limit ASA's hydrolysis, which produces a product interfering with sizing.

[0006] Methods have been proposed to prevent ASA's hydrolysis. For example, a water emulsion containing cationic starch is prepared as late as possible before application. However, the process for mixing the ASA with cationic starch is rather complicated and uses a complex machine.

[0007] Another known hydrophobation method used to increase paper water resistance is vapor depositing the ASA on the paper surface. In this case, ASA in gaseous phase contacts the paper surface and reacts with the hydroxyl groups of the cellulose. However, applying such a method at an industrial scale would imply using a complex system in order to avoid releasing ASA in the atmosphere surrounding the machine. A complicated system would be required for confining the molecules in gaseous phase while allowing their contact with the paper sheet in continuous movement.

[0008] In light of the aforementioned, there is thus a need for new ASA containing sizing compositions as an alternative to ASA water based compositions.

SUMMARY

[0009] It is therefore an aim of the present invention to address the above mentioned issues.

[0010] In one aspect, there is provided a water-free surface sizing composition comprising at least one alkenyl succinic anhydride and at least one biosolvent for adjusting a viscosity of the composition to allow spraying thereof on a paper surface.

[0011] In another aspect, there is provided a method for treating a paper substrate with the water-free surface sizing composition, comprising spraying the composition onto the surface of the paper substrate and heating the paper substrate treated with the composition.

[0012] In another aspect, there is provided a use of the water-free surface sizing composition for providing water resistance or improving water resistance of a paper substrate.

[0013] Other objects, advantages and features of the present invention will become more apparent upon reading of the following non-restrictive description of embodiments thereof.

DETAILED DESCRIPTION

[0014] A water-free sizing composition providing water-resistance to the surface of paper substrates will be described.

[0015] In the present description, the expression “paper substrate” refers to any type of cellulosic fiber-based substrate including, for example, and without being limited to, any suitable wood-fiber based material, such as recycled or virgin liner, medium, chipboard, paperboard, folding carton, kraftpak paper, bag paper, fine paper and the like.

[0016] The cellulose in the cellulosic fiber-based substrate is accessible for surface treatment. In other words, hydroxyl groups of the cellulose included in the paper substrate are accessible for reacting with the ASA molecules of the water-free composition. If the cellulose in the paper substrate has previously been surface treated, for example, with starch, remaining hydroxyl groups have to be accessible for reacting with the ASA of the water-free composition. In one embodiment, the cellulose in the paper substrate to be treated with the water-free sizing composition containing ASA has not been subjected to a previous surface treatment. In another embodiment, the paper substrate may have been subjected to an internal sizing treatment prior to the surface treatment with the water-free composition. The internal sizing treatment may be carried out using any sizing agent known in the art for internal treatment. For example, the internal sizing agent can be an AKD or an ASA. When an ASA is used as internal sizing agent, it can be the same or different than the ASA present in the water-free composition with biosolvent.

[0017] Broadly described, the sizing composition is a water-free or substantially water-free solution of at least one ASA in a biosolvent. The composition has a viscosity that allows it to be sprayed on the paper substrate.

[0018] By “sprayed” or “spraying”, it is meant that the composition is applied as a liquid broken up into minute droplets being blown, ejected into, or falling through the air to then reach the surface of the paper substrate. In an embodiment, the composition is applied to the paper surface in the liquid state either using a shower or by discharge from a pressurized container through spray nozzles.

[0019] ASAs are liquid products having a relatively high viscosity. Mixing the ASA with a biosolvent or mixture of biosolvents preferably reduces the viscosity of the ASA in the resulting composition. With an appropriate viscosity, the composition can be applied by spraying on the paper substrate. This results in a substantially homogeneous distribution of the ASA onto the paper surface. The ASA is thus allowed to react with accessible hydroxyl groups of the cellulose in the paper substrate and hydrophobicity of the paper substrate is thus increased.

[0020] The ASAs that can be used in the composition include any ASA commonly used as an internal sizing agent in the paper industry. It is also possible to use a mixture of different ASAs in the composition.

[0021] In an embodiment, the ASA has an alkenyl group of from 16 to 20 carbon atoms. In another embodiment the ASA has an alkenyl group of from 16 to 18 carbon atoms. When the composition contains a mixture of ASAs wherein each has from 16 to 20 carbon atoms in its alkenyl group. Moreover, the double bond of the alkenyl group can be in any position on the alkenyl chain.

[0022] According to another embodiment, the ASA used in the composition include hexadecenyl succinic anhydride, octadecenyl succinic anhydride or any mixture thereof, wherein the double bond of the alkenyl group is in any position on the alkenyl chain.

[0023] In another embodiment, the ASA added to the composition is present in the product NALSIZE® 7542, sold by Nalco Company, or HYDRORES™ AS 2300, sold by Kemira Chemicals. NALSIZE 7542 is a mixture of ASAs (C₁₆–C₁₈) containing up to 2% nonionic surfactant. HYDRORES AS 2300 is ASA having a linear alkenyl chain of 18 carbon atoms.

[0024] In the composition, the ASA or mixture of ASAs are combined with at least one biosolvent to decrease the viscosity of the ASA(s). In some embodiments, a mixture of biosolvents can be used to achieve the required viscosity. The mixture ASA(s)- biosolvent(s) is a liquid solution that is substantially homogeneous.

[0025] Biosolvents, as opposed to petroleum-derived solvents, are solvents from natural origin which are issued from treated or untreated plant, animal or mineral raw materials.

[0026] Examples of biosolvents which can be used in the composition include biodiesels which are vegetable oil- or animal fat-based diesel fuel comprising long-chain alkyl (*e.g.*, methyl, propyl or ethyl) esters.

[0027] Other examples of biosolvents include dipentene, the racemic of (+) and (-) limonene. It is also possible to use only one of the enantiomers of limonene.

[0028] Other possible biosolvents to be used in the composition include fatty acid esters and fatty acid amides. The fatty acid esters or amides are either saturated or unsaturated. In an embodiment, the fatty acid esters are fatty acid methyl esters and the fatty acid amides are N,N-dimethyl fatty acid amides. In some embodiments, the aliphatic chain of the fatty acid esters has from 8 to 18 carbon atoms. Examples of fatty acid esters include methyl caprylate, methyl laurate, methyl olete, or methyl palmitate. In some embodiments, the aliphatic chain of the fatty acid amides has 8 or 10 carbon atoms. The fatty acid amides may be N,N-dimethylcaprylamide or N,N-dimethylcapramide.

[0029] As previously mentioned, the use of a biosolvent or a mixture of biosolvents can be utilized to decrease the viscosity of the ASAs, thereby obtaining a sizing composition that is sprayable. In an embodiment, the biosolvent and its relative amount in the composition are determined to achieve a composition with a viscosity of about 100 cPs or less. In another embodiment, the composition has a viscosity of from about 25 to about 100 cPs. In some embodiments, the viscosity of the composition can be between about 25 and about 90 cPs.

[0030] The term “about” as used in the present description means within an acceptable error range for the particular value as determined by one of ordinary skill in the art, and will depend in part on how the value is measured or determined, *i.e.*, the limitations of the measurement system. It is commonly accepted that a 10% precision measure is acceptable and encompasses the term “about.”

[0031] In another embodiment, the sizing composition has a flash point of at least about 50°C. The value of the flash point of the composition will principally depend on the nature and proportions of the biosolvent(s) used in the composition. The proportion of ASAs is also taken into account. ASAs have high flash points and contribute to an increase of the flash point of the composition. A person skilled in the art will be able to choose the appropriate biosolvents and to estimate the proportions thereof to obtain a composition with an appropriate flash point. In some embodiments, the flash point is chosen so as to minimize

flammability risks of the composition in the dryer or through the contact with hot surfaces during the sizing process. For example, the flash point of the composition may be of at least about 93°C.

[0032] The surface sizing composition is obtained by mixing the ASA or mixture of ASAs with the biosolvent or mixture of biosolvents.

[0033] In an embodiment, the ASA or mixture of ASAs are added in about 1 wt% to about 80 wt% of the weight of the composition. In another embodiment, ASA(s) represent(s) about 40 wt% to about 70 wt% of the weight of the composition.

[0034] The biosolvent or mixture of biosolvents can be present in about 20 wt% to about 99 wt% of the weight of the composition, or in about 30 wt% to about 60 wt% of the weight of the composition.

[0035] In another embodiment, the surface sizing composition comprises about 60% w/w of ASA and about 40% w/w of biosolvent or mixture of biosolvents. The ASA can be NALSIZE 7542 and the biosolvent a mixture of biodiesel and limonene.

[0036] Table 1 below provides examples of sizing compositions according to specific embodiments.

Table 1:

	Composition 1	Composition 2
NALSIZE 7542	60% w/w	60% w/w
Biodiesel	38% w/w	35% w/w
Limonene	2% w/w	5% w/w
Viscosity 50 rpm, spindle #1	43 cp	41 cp
Flash point	107°C	95°C

[0037] An embodiment of a method for treating a paper substrate with the sizing composition will be now described.

[0038] The method generally involves spraying the composition onto the surface of the paper substrate and then heating the treated substrate.

[0039] The paper substrate, which can be a recycled or virgin liner, medium, chipboard, folding carton, kraftpak paper, paperboard, bag paper, fine paper or any other cellulosic fiber-based substrate, is provided to the sizing machine where the composition is allowed to be sprayed on its surface using a sprayer.

[0040] In an embodiment, the composition is applied to the paper surface in the liquid state using a shower or any spray equipment commonly known in the art. For example, the composition can be applied by discharge from a pressurized container through a multi-nozzles spraying system. Alternatively, the composition can be applied using a rotor damping system, for instance a WEKO-RFT Rotor Damping System. When the water-free composition is sprayed using a multi-nozzles spraying system, the nozzles can be appropriately placed across the width of the paper machine. The spray nozzles are designed and spaced to ensure even distribution of the composition on the paper sheet.

[0041] In an embodiment, the composition is applied at room temperature on the paper surface. The quantity of composition applied to the surface of the paper substrate may depend on the type of substrate and the intended water barrier. In an embodiment, the quantity of composition applied to the surface of the paper substrate is from about 0.2 to about 10 g/m². In another embodiment, the quantity of composition applied to the surface of the paper substrate is from about 0.2 to about 2 g/m².

[0042] Once the water-free sizing composition has been applied on the surface of the paper, the treated paper is then passed through a dryer or heater to provide the energy required to allow the reaction between the hydroxyl groups of the cellulose included in the paper and the ASA molecules, and the surface of the substrate becomes hydrophobic.

[0043] Usually, dryers/heaters commonly used in paper making processes are adapted for heating the paper treated with the water-free composition and there is no need to modify their temperature.

[0044] As previously mentioned, the water-free surface sizing composition once applied to the paper and after heating thereof, provides good water resistance properties to the paper. The so treated paper can show Cobb_{2min} values from about 27 g_{water}/m² to about 50 g_{water}/m². The so treated paper can be used in many applications, for example, printing paper, linerboard, for folding box and protective headers.

[0045] The present water-free sizing composition and the way it is applied to the paper substrate show various advantages over known paper sizing methods. The use of a water-free sizing composition allows avoiding paper curl problems that can be observed when applying water based sizing compositions using showers.

[0046] The present water-free composition, thanks to the biosolvents it contains, is more environmentally friendly than compositions containing petroleum based solvents.

[0047] The following examples are provided to illustrate some properties and advantages of the coating.

EXAMPLES

Example 1:

[0048] Water-free surface sizing compositions have been prepared as summarized in Table 2. Their viscosities and flash points have been determined and are also reported in Table 2.

Table 2 :

Composition		B1	B2	B3	T1	T2	T3	T4	T6	D3	D5	T12	T13
ASA*	(w/w %)	1	1	1	60	60	60	60	60	75	90	60	80
Biodiesel**	(w/w %)	99	0	0	40	35	30	20	0	0	0	38	18
Limonene	(w/w %)	0	0	99	0	5	10	20	40	25	10	2	2
Dipentene	(w/w %)	0	99	0	0	0	0	0	0	0	0	0	0
Viscosity, spindle #1, 22°C	(cP)	23	11	10	44	41	38	32	24	40	80	43	85
Flash point	(°C)	120	50	55	180	95	83	67	54	65	78	107	109

* ASA is NALSIZE 7542.

** Biodiesel derived from vegetable oils

Example 2:

[0049] Compositions B1, B2 and B3 of Table 1 were tested to evaluate their sizing properties.

[0050] 2.4 g paper handsheets were prepared using brown pulp (100% old corrugated containers ("OCC")). The retention system was composed of 0.6 kg/t PERCOL® 3320 CB ("C-PAM") (polyacrylamide, available from BASF) and 4 kg/t LUREDUR® 8097 (partially hydrolyzed polyvinyl formamide, available from BASF). The paper handsheets dryness before applying the compositions was 34%.

[0051] The compositions were applied onto the surface of the 2.4 g paper handsheets using an aerograph. The liquid compositions were uniformly vaporized using compressed air. The handsheets were then dried at 105°C for 15 minutes and left for 5 days at 23°C under

50% relative humidity. Cobb_{2 min} values were then measured. The results are reported in Table 3. Measurements were also performed for an untreated paper handsheet for comparison. The Cobb_{2 min} for the untreated substrate was above 220 g_{water} / m².

Table 3:

Solution		B1	B2	B3
ASA	(%)	1	1	1
Biodiesel	(%)	99	0	0
Limonene	(%)	0	0	99
Dipentene	(%)	0	99	0
Cobb _{2 min}	(g _{water} / m ²)	36	42	27

[0052] The results presented in Table 3 show that the paper substrate was successfully treated with compositions B1–B3. The paper substrate treated with any one of compositions B1–B3 has an improved water resistance compared to the untreated substrate, even with a very low content of the ASA in the composition.

Example 3:

[0053] A composition was prepared by mixing 60 % (w/w) NALSIZE 7542 as ASA, and a mixture of 35 % (w/w) biodiesel and 5 % (w/w) limonene as biosolvent. The composition was applied to the surface of a cardboard (recycled paper; basis weight 679 g/m²), at the mill before the dyer section, using a spray gun. The sizing efficacy was studied over time by measuring Cobb values four times within a period of one year and 4 months. The treated cardboard was not oven-dried. The untreated surface allowed water penetration into the cardboard on the Cobb_{2 min} test (about 967 g_{water} / m²).

[0054] The results of the Cobb tests are provided in Table 4.

Table 4:

Time after treatment	Cobb _{2 min} (g _{water} / m ²)	Cobb _{15 min} (g _{water} / m ²)	Cobb _{30 min} (g _{water} / m ²)
0 *	31	70	n.d.
2 weeks	32	71	n.d.

4 weeks	30	70	n.d.
1 year and 4 months	n.d.	70	102

* measurements were performed on the cardboard immediately after treatment

n.d.: not determined

[0055] The values of $\text{Cobb}_{2 \text{ min}}$ (about $30 \text{ g}_{\text{water}}/\text{m}^2$) and $\text{Cobb}_{15 \text{ min}}$ (about $70 \text{ g}_{\text{water}}/\text{m}^2$) show that the application of the ASA in biosolvents increases the hydrophobicity of the cardboard surface compared to the untreated cardboard ($\text{Cobb}_{2 \text{ min}}$ of about $967 \text{ g}_{\text{water}}/\text{m}^2$).

[0056] The above-described embodiments and examples are considered in all respect only as illustrative and not restrictive, and the present application is intended to cover any adaptations or variations thereof, as apparent to a person skilled in the art. Of course, numerous other modifications could be made to the above-described embodiments without departing from the scope of the invention, as apparent to a person skilled in the art.

[0057] The scope of the invention is therefore intended to be limited solely by the scope of the appended claims.

CLAIMS

We claim:

1. A water-free surface sizing composition comprising at least one alkenyl succinic anhydride and at least one biosolvent.
2. The composition of claim 1, wherein the composition has a viscosity of from about 25 cPs to about 100 cPs.
3. The composition of any of claims 1 to 2, wherein the at least one alkenyl succinic anhydride comprises an alkenyl succinic anhydride having an alkenyl group of from 16 to 20 carbon atoms.
4. The composition of any of claims 1 to 3, wherein the at least one alkenyl succinic anhydride comprises an alkenyl succinic anhydride selected from the group consisting of hexadecenyl succinic anhydride, octadecenyl succinic anhydride, and combinations thereof.
5. The composition of any of claims 1 to 4, wherein the at least one biosolvent comprises at least one of biodiesel, dipentene, limonene, a fatty acid ester, and a fatty acid amide.
6. The composition of claim 5, wherein the at least one biosolvent comprises a fatty acid ester.
7. The composition of claim 6, wherein the fatty acid ester has an aliphatic chain of from 8 to 18 carbon atoms.
8. The composition of claim 6, wherein the fatty acid ester is selected from the group consisting of methyl caprylate, methyl laurate, methyl olate, methyl palmitate, and combinations thereof.
9. The composition of claim 5, wherein the at least one biosolvent comprises a fatty acid amide.
10. The composition of claim 9, wherein the fatty acid amide is an N,N-dimethyl fatty acid amide.

11. The composition of claim 10, wherein the N,N-dimethyl fatty acid amide has an aliphatic chain of at least one of 8 and 10 carbon atoms.
12. The composition of claim 9, wherein the fatty acid amide is at least one of dimethylcaprylamide and N,N-dimethylcapramide.
13. The composition of any of claims 1 to 12, wherein the at least one alkenyl succinic anhydride is present in the composition at a concentration of from about 1 wt% to about 80 wt%.
14. The composition of any of claims 1 to 13, wherein the at least one biosolvent is present in the composition at a concentration of from about 20 wt% to about 99 wt%.
15. The composition of claim 1, wherein the at least one alkenyl succinic anhydride comprises an alkenyl succinic anhydride having an alkenyl group of from 16 to 20 carbon atoms and is present in the composition at a concentration of from about 1 wt% to about 80 wt%, wherein the at least one biosolvent comprises at least one of biodiesel, dipentene, limonene, a fatty acid ester, and a fatty acid amide and is present in the composition at a concentration of from about 20 wt% to about 99 wt%, and wherein the composition has a viscosity of from about 25 cPs to about 100 cPs.
16. The composition of claim 1, wherein the at least one alkenyl succinic anhydride comprises an alkenyl succinic anhydride having an alkenyl group of from 16 to 20 carbon atoms and is present in the composition at a concentration of from about 40 wt% to about 70 wt%, wherein the at least one biosolvent comprises at least one of biodiesel, limonene and dipentene and is present in the composition at a concentration of from about 20 wt% to about 99 wt%, and wherein the composition has a viscosity of from about 25 cPs to about 100 cPs.
17. A method of treating a paper substrate having a surface, the method comprising:
 - spraying the composition of any of claims 1 to 16 onto the surface of the paper substrate, thereby forming a treated substrate; and
 - heating the treated substrate.
18. The method of claim 17, wherein the paper substrate was subjected to an internal sizing treatment prior to the spraying.

19. The method of claim 18, wherein the internal sizing treatment comprises applying at least one of alkyl ketene dimer and alkeny succinic anhydride.

20. The method of any of claims 17 to 19, wherein the composition is sprayed at from about 0.2 gram to about 10 grams per square meter of surface of the paper substrate.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2014/042293**A. CLASSIFICATION OF SUBJECT MATTER****D21H 21/16(2006.01)i**

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)

D21H 21/16; C08H 500; D21H 19/76; D21H 17/72; D21H 17/04; D21H 17/06; D21H 17/14; C07G 1700

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

Korean utility models and applications for utility models

Japanese utility models and applications for utility models

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

eKOMPASS(KIPO internal) & Keywords:paper, cellulose, sizing, water free, alkenyl succinic anhydride, ASA

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 01-44575 A1 (AKZO NOBEL N.V. et al.) 21 June 2001 See abstract; page 8, lines 16-18; and claims 1, 10.	1-3, 15-16
Y	US 6273997 B1 (EHRHARDT, S. M. et al.) 14 August 2001 See abstract; and claims 1, 9.	1-3, 15-16
A	US 6692560 B2 (HOLMBERG, K. et al.) 17 February 2004 See abstract; and claims 1, 11.	1-3, 15-16
A	US 6485555 B1 (LINDGREN, E. et al.) 26 November 2002 See abstract; and claim 1.	1-3, 15-16
A	WO 2005-118953 A1 (FUJI PHOTO FILM B.V.) 15 December 2005 See abstract; and claim 1.	1-3, 15-16



Further documents are listed in the continuation of Box C.



See patent family annex.

* 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 but 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

10 October 2014 (10.10.2014)

Date of mailing of the international search report

10 October 2014 (10.10.2014)

Name and mailing address of the ISA/KR

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INTERNATIONAL SEARCH REPORTInternational application No.
PCT/US2014/042293**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.: 6-12, 18-19
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:
Claims 6-12 and 18-19 are unclear because they substantially refer to multiple dependent claims 5 and 17 which do not comply with PCT Rule 6.4(a).
3. ☒ Claims Nos.: 4-5, 13-14, 17, 20
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 an 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

Information on patent family members

International application No.

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