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Publication number:

**0 177 113
B1**

12

EUROPEAN PATENT SPECIFICATION

45 Date of publication of patent specification: 19.07.89

51 Int. Cl.⁴: D 21 C 9/06, D 21 C 9/08

21 Application number: 85302568.2

22 Date of filing: 12.04.85

54 Improved method of brown stock washing.

38 Priority: 19.09.84 US 652311

43 Date of publication of application:
09.04.86 Bulletin 86/15

45 Publication of the grant of the patent:
19.07.89 Bulletin 89/29

84 Designated Contracting States:
DE FR GB SE

56 References cited:
US-A-1 730 387
US-A-2 144 756
US-A-3 992 249
US-A-4 184 912
US-A-4 297 164

ABSTRACT BULLETIN OF THE INSTITUTE OF
PAPER CHEMISTRY, vol. 54, no. 8, February
1984, page 985, no. 9251, Appleton, Wisconsin,
US ; & JP - A - 83 13 795 (NIPPON OILS & FATS
CO., LTD.; KURITA INDUSTRY CO., LTD.)
26-01-1983

TAPPI, vol. 63, no. 2, February 1980, pages
81-87, Atlanta, GA, US; L.H. ALLEN:
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newsprint mills"

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Courier Press, Leamington Spa, England.

EP 0 177 113 B1

Description**Field of the Invention**

This invention relates to a method of removing and recovering spent or excess cooking chemicals and pitch from virgin or primary cellulosic fiber. More particularly, the invention relates to an improved method of brown stock washing to enhance the recovery of spent cooking chemical residues and excess cooking chemicals, and remove non-cellulosic materials such as lignin, fatty acid soaps, and resin acids.

Background of the Invention

Virgin cellulosic fiber typically derived from logs of hardwood or softwood, undergoes lengthy processing before it is suitable for use in papermaking. In a typical pulping process, briefly, logs are reduced to wood chips, which are fed into a digester. "Liquor", an aqueous solution obtained from the later described wash step and containing dissolved and residual cooking chemicals, spent cooking chemical residue and cellulosic contaminants, and "white liquor", another by-product of the pulping process known in the art, are fed into the digester, primarily for dilution. Cooking chemicals are also added as required. The cooling chemicals are described hereinafter.

The contents of the digester are brought to a relatively high temperature and pressure, for example about 177°C (about 350°F) at a pressure of about 760000 N/m² (about 110 pounds per square inch). The wood chips are "cooked" in the digester under these conditions to reduce the wood chips to pulp. Typically, under these conditions, the wood chips are cooked from about 1 to 5 hours. The cooking can be carried out in batch or continuous digesters.

The cooked wood chips or pulp in the aqueous medium after digestion is referred to as "brown stock". The brown stock consists generally of two phases, the pulp, and the liquor or liquid phase of the digester contents. However, typically after digesting, oversized chips, insufficiently cooked chips, or knots remain. These components are generally removed from the brown stock by knotters which typically consist of coarse screens.

Before further processing of the pulp, it is generally considered necessary to separate the pulp from the liquor. It is also desirable to clean the pulp, removing and to the greatest extent possible, recovering spent or excess cooking chemicals, and removing and recovering pitch contaminants.

After digestion, and following removal of oversized chips and the like, the brown stock is transferred to a washer for a washing step. Typically, the washing process involves a series of washers which separate the pulp from the liquor, and progressively clean the pulp by removal of cooking chemicals, cooking chemical residues, and non-cellulosic contaminants.

Several methods may be used to perform the washing step. In the past, the brown stock was filtered in a false bottom tank or diffuser into which the digester was discharged. The liquor was drained through the false bottom, and the pulp was washed by gravity displacement of the liquor with wash water. Other types of washers such as a pressure washer are also known in the art.

Currently, the rotary vacuum drum or cylinder or vacuum washer is more typically used. As is known to those familiar with the art, the vacuum washer is generally a wire cylinder or drum that rotates in a vat containing the brown stock (i.e. the pulp and liquor mixture). The lower section of the drum is immersed in the brown stock. Vacuum is applied inside the drum as it rotates through the brown stock. The liquor drains through the surface of the wire drum into the interior, leaving a layer of pulp on the outside face of the drum. The layer of pulp is held in place by the vacuum force inside the drum, from where it is conducted away.

The layer of pulp continues to build, forming a mat or sheet, as the submerged portion of the drum rotates through the brown stock in the vat. Liquor continues to drain from the pulp of fiber mat as a result of the differential pressure between the external atmosphere and the vacuum within the cylinder.

Washing action is generally provided by showers located over the pulp sheet. Water is sprayed onto the pulp sheet to displace the liquor from the sheet on the drum as the drum continues to rotate. The vacuum force draws the water into the sheet, where it displaces the liquor. The liquor drains out the other side of the sheet into the inside of the cylinder, where it drains away to a filtrate storage tank for reuse, for example as wash water for a more contaminated sheet which has formed on another of the washers in the series.

Finally, the pulp sheet is removed from the face of the wire by a doctor blade.

The surface of the sheet where the wash water is applied becomes cleaner than the pulp adjacent to the cylinder at the bottom of the sheet, since the wash water becomes more concentrated in liquor as it passes through the sheet. Consequently, where a series of washers is utilized, the pulp sheet obtained from the first vacuum washer is generally repulped to provide a more uniformly clean pulp before traveling over the second vacuum washer. This repulping step is generally repeated between each vacuum washer in the sequence.

In the repulping step, the pulp fibers are agitated at a low consistency (i.e. the pulp is very dilute) in order to facilitate scrubbing. The low consistency also aids in achieving a lowered concentration of dissolved solids, prior to collection of the pulp on the next washer in the series. Low consistency promotes diffusion of the contaminated liquor from the pulp in the repulping step.

In a sequence of washers, the pulp medium and the wash water are generally arranged to flow

countercurrent to each other. Fresh water is typically used to wash the pulp sheet on the last stage washer. The filtrate that was pulled through pulp sheet on each washer is used to wash the pulp on the preceding washer. This aids in minimizing dilution of the liquor which is separated from the pulp, and from which cooking chemicals or cooking chemical residues are to be recovered as described hereinafter.

5 The cooking chemicals used in pulping mills are known in the art. Briefly, the cooking system is generally either kraft or sulfite. Other cooking systems are also known in the art.

The kraft system generally involves the use of sodium hydroxide and sodium sulfide in the digester to aid in decomposition of the wood fibers to produce pulp. The sodium may be added as sodium sulfate, sodium carbonate, or similar sodium compounds. The sulfite system typically involves the use of SO₂ and magnesium, calcium, sodium or ammonia. The kraft mill generates "black liquor", while the counterpart in the sulfite mills is referred to as "red liquor". For the purpose of this description, the term "liquor" refers to both "red" and "black" liquor, and the aqueous phase of the pulp mixture resulting from other pulp processing methods such as those described below.

15 Some pulping mills form pulp from wood products without the use of cooking chemicals. Several such pulping processes are known, including mechanical processes such as the groundwood process, use of a refiner to create refiner mechanical pulp, or use of heat to create thermomechanical pulp. Most such processes rely on heat and mechanical action to break down the wood fibers. Other processes, such as the NSSC process, rely on both chemical and mechanical action. While these mechanical, thermomechanical, or semi-chemical processes typically do not involve washing steps, where washing steps are used the methods of this invention can aid in cleaning the pulp and recovering organic contaminants.

20 Before washing, the brown stock will contain many impurities from the pulping process, including excess cooking chemicals and spent cooking chemicals (where chemicals are used in the pulping), and also a variety of organic contaminants such as resin acids, fatty acid soaps and the like originating in wood. The contaminants occlude to the pulp fibers, and are also present in the aqueous phase of the brown stock. It has been found that in general, the contaminants of black liquor and the corresponding pulp are principally alkali lignin, hydroxy acids and lactones, and sodium. Generally, black liquor is also contaminated with acetic acid, formic acid, sulfur, extractives, and methanol. Red liquor (obtained through the sulfite process) and the corresponding pulp has been found to be contaminated with lignosulfonate, monosaccharides (mannose, xylose, galactose, glucose and arabinose), poly and oligosaccharides, calcium, aldonic acids, sugar-sulfonates, extractives, acetic acid, methanol, and glucuronic acid. These materials are substantially different from those encountered in deinking or dewaxing repulping processes, where the contaminants are generally inorganic substances and very different organic compounds.

25 It is highly desirable to recover or reclaim the cooking chemical residues for reuse, to reduce the amount of chemicals which must be purchased by the mill. Those residues which remain in the pulp after the pulping process are generally not recovered, and contaminate the pulp products. Those residues which are carried by the liquor tend to be recoverable. Therefore, it is advantageous to decrease the amount of chemicals carried by or occluded to the pulp and increase the amount carried by the liquor. In particular, it is desirable to effect a transfer of chemicals from the pulp to the liquor.

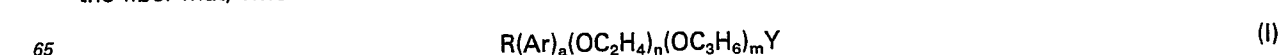
30 Such a transfer can be achieved to a great extent by washing of the pulp. However, for production level washing to obtain large quantities of pulp at a high quality level or level of purity, vast quantities of wash water are required. The wash water dilutes the liquor and chemicals washed from the pulp. Since recovery of the chemicals involves distillation, or evaporation of the aqueous component, it can be significantly more expensive to recover chemicals from a more dilute solution, off-setting the cost benefits to be achieved by recovery. Thus, a substantial need exists for a method of washing virgin pulp which will sufficiently remove excess and spent cooking chemicals from the pulp and otherwise clean the pulp without causing excessive dilution.

35 In addition to the problems of recovering inorganic cooking chemicals residues, another similar set of second problems encountered in pulping virgin pulp results from the presence of lignin and other organic substances such as resin acids, fatty acid soaps, etc. in wood chips. It is desirable to recover these substances because they are economically or commercially valuable, for example, when recovered as tall oils. In addition, pulp which retains a high level of such materials may require the use of more chemicals in the bleaching step, thus rendering the bleaching step more costly. A need exists for a method of washing virgin pulp which will result in greater recovery of organic substances and lighter colored pulp while minimizing the amount of dilution or wash water required to obtain these results.

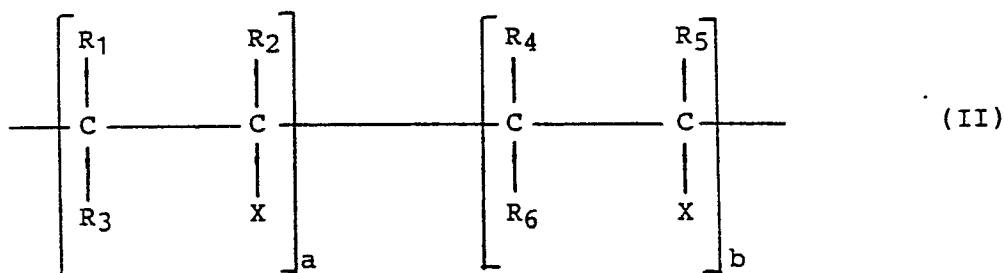
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Brief Description of the Invention

According to the present invention, there is provided an improved method of removing spent or excess cooking chemical compounds or organic contaminants from chemically or mechanically prepared virgin pulp, characterised in that it comprises (a) forming a fiber mat of chemically or mechanically prepared virgin pulp; and (b) forcing an aqueous solution comprising a substituted oxyethylene glycol non-ionic surfactant, a water soluble low molecular weight polyelectrolyte dispersant and a solvent into and through the fiber mat, wherein the non-ionic surfactant has the formula:



wherein Ar is an aromatic residue, a is zero or 1, R is an aliphatic group that contains from 6 to 24 carbon atoms when a is zero and from 6 to 18 carbon atoms when a is 1, n is from 3 to 50, m is from zero to 50, and Y is hydroxy or benzyl ether with the proviso that, when m is zero, Y is benzyl ether, and wherein the polyelectrolyte dispersant has the formula:



wherein R₁, R₂, R₄ and R₅ are independent and are selected from hydrogen, C₁—C₄ lower alkyl, carboxy-substituted alkyl or mixtures thereof, R₃ and R₆ are independent and selected from hydrogen, carboxy, carboxy-substituted alkyl or mixtures thereof, X is selected from carboxy, salts and derivatives of carboxy, acetyl, hydrocarbon moieties commonly attached to free radical monomers, COOZ where Z is H, a monovalent metal ion or ammonium, or mixtures thereof, and the total of a + b falls in the range of 15 to 1,000.

The present invention also provides a method of recovering excess or spent cooking chemical compounds or organic contaminants from an aqueous pulping medium comprising chemically or mechanically prepared virgin pulp and excess or spent chemical compounds or organic contaminants, characterised in that it comprises (a) combining the aqueous pulping medium with a substituted oxyethylene glycol non-ionic surfactant, a water soluble molecular weight polyelectrolyte dispersant, and solvent; (b) substantially separating the chemically or mechanically prepared virgin pulp from the aqueous pulping medium; and (c) after step (b), removing excess or spent cooking chemical compounds or organic contaminants from the aqueous pulping medium; wherein the non-ionic surfactant and the polyelectrolyte dispersant have the formula (I) and (II) given above.

We have found that the addition of the nonionic surfactant, the dispersant, and the solvent to the wash water or the brown stock itself will result in unexpectedly improved washing of virgin pulp. By the method of this invention, a given quantity of wash water will result in surprisingly increased removal of spent or excess cooking chemical compounds and organic contaminants, thus minimizing dilution in the washing process while providing cleaner, and generally lighter colored, pulp and facilitating economical recovery of the cooking chemical residues and organics.

Detailed Description of the Invention

The present invention involves use in the wash process of the nonionic surface active agent or surfactant in combination with the dispersant and the solvent. The surface has the formula (I) given above wherein Ar is preferably monocyclic and R is typically saturated.

The foregoing structural formula (I) can be considered to encompass two major classes of surfactants, i.e. (a) alkylene oxide adducts of alkylphenols, and (b) alkylene oxide adducts of higher (greater than C₅) aliphatic alcohols or acids. Acids which can be utilized in the formation of the surfactant include lauric, myristic, oleic, linolenic, palmitic and stearic. Where the adduct of an aliphatic acid is used, may contain some unsaturation.

The surfactants contemplated for use in the invention are generally low foaming surfactants which do not significantly contribute to foam problems within the system.

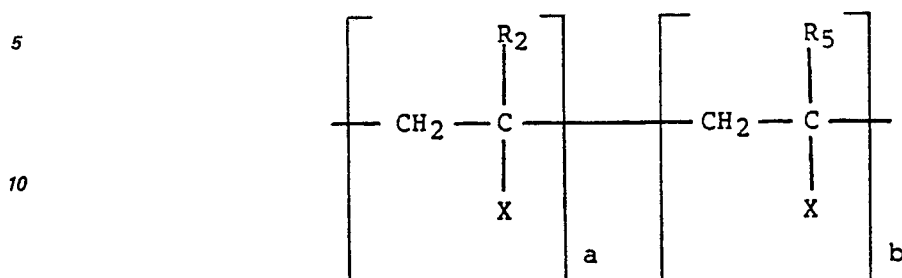
These surfactants are described in detail in commonly assigned United States Patent Application Serial No. 093,744 filed November 13, 1979 on behalf of Richard E. Freis, James E. Maloney and Thomas R. Oakes, entitled "Methods of Deinking Secondary Fibers", the entire disclosure of which (and of its equivalent GB—A—2063326) is incorporated by reference herein. A continuation of U.S. Serial No. 093,744 was filed on January 17, 1983, and has been assigned U.S. Serial No. 458,432.

The present invention also involves the use of a water soluble low molecular weight polyelectrolyte dispersants having the formula (II) given above, which covers any homo, co, ter, etc., polymer of the formula (II). An example of the carboxy-substituted alkyl group in formula (II) is —CH₂COOH, an example of the derivative of a carboxy group in formula (II) is amide, and an example of the hydrocarbon moiety commonly attached to free radical polymerizable monomers in formula (II) is —C₆H₅ in styrene.

Examples of materials within the scope of the above formula (II) include polymaleic acid, polyacrylic acid, polymethacrylic acid, polyacrylic acid/itaconic acid copolymers, polyacrylic acid/hydrolyzed maleic acid copolymers, polymaleic acid/itaconic acid copolymers, hydrolyzed polymaleic acid/vinyl acetate copolymers, polyacrylic acid/acrylamide copolymers, polyacrylic acid/methacrylic acid copolymers, styrene/maleic acid copolymers, sulfonated styrene/maleic acid copolymers, polymaleic acid/methacrylic acid copolymers, maleic acid telomers, maleic/alkyl sulfonic copolymers.

EP 0 177 113 B1

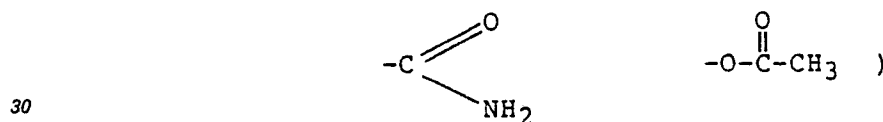
A particularly preferred class of water soluble polyelectrolytes for use in the practices of the present invention is the polyacrylate compounds. The polyacrylate compounds comprise polymers and copolymers of the structure:



and their derivatives, wherein R_2 , R_5 , X , a and b are defined as above.

In a most preferred practice of the present invention, X is $-\text{COOZ}$, wherein Z is H , or a monovalent cation, e.g. Na^+ , K^+ , or NH_4^+ . Thus, typical of the preferred polyelectrolytes of the present invention are polyacrylic acid, polymethacrylic acid and copolymers of acrylic acid/methacrylic acid (e.g., AQUATREAT available from ALCO Chemical). The words aquatreat and ALCO, and the words CELLOSOLVE and CARBITOL used below, are Registered Trade Marks.

The polyelectrolytes used in this invention must be water soluble. Generally speaking, to be water soluble, the polymer must contain sufficient polar groups (e.g., COOH) for the molecule to interact with the polar water molecules. This means that in copolymers, terpolymers, tetramers, etc., with unsaturated monomers which are predominantly or entirely hydrocarbon (e.g., styrene) there must be sufficient polar functional groups for the polymer to dissolve in room temperature or below water. Generally, at least about 10 mole percent of the monomers comprising the polymer must contain polar functionality (e.g.,



to provide the required water solubility.

The low molecular weight polyelectrolytes used in present invention generally have molecular weights of less than about 50,000 with preferred molecular weights in the range of about 500 to 25,000, most preferably of 750 to 5,000. Thus, the sum of $a + b$ above, generally falls in range of 5 to 1,000, preferably 10 to 500 and most preferably 12 to 450. One skilled in the art will recognize that the materials within the above molecular weight ranges are generally of lower molecular weight than polymers generally referred to in the art as flocculants which may have molecular weights in the range of several million or more. Flocculants perform function of agglomerating suspended particles opposite the desired function of dispersion described herein. Thus, these high molecular weight materials operate in a manner effectively opposite that of the materials described herein. The lower molecular weight materials of the present invention are generally referred to in the art as "dispersants".

The improvement of the present invention contemplates the use of various well-known water soluble solvents or cosolvents, along with the dispersants and surfactants. The solvents unexpectedly provide increased removal of contaminants of the pulp. The solvents can be ethoxylated solvents such as the glycol ethers available under the trademarks Cellosolve and Carbitol. Preferred examples of solvents for use in this invention include tetrahydrofuran, tetrahydrofurfuryl alcohol, and ethoxylated and propoxylated derivatives thereof. It has been found that tetrahydrofurfuryl alcohol is particularly beneficial in the context of the invention, and it is theorized that this component contributes to the high recovery of pulp purity obtained by the method of this invention.

Functionally speaking, the nonionic surfactant, dispersant, and solvent additives of the invention should be utilized in sufficient amounts of ratios to achieve increased recovery of cooking chemicals and soluble organics, and increased pulp purity after washing with a given volume of water. We have found that the components produce the best results at a surfactant:dispersant ratio from about 0.5:1 to 2:1. We have found the more effective ratios of surfactant:solvent to be from about 0.5:1 to 1:1.

As used in brown stock washing, we have found the desired concentration of the nonionic surfactants in the context of this invention to be generally in the range of about 0.005 to 15 kg/metric ton (about 0.01 to 30 lbs/short ton) oven dried pulp, with concentrations in the range of about 0.075 to 2.5 kg/metric ton (about 0.15 to 5 lbs/short ton) being preferred. The concentrations of the polyelectrolyte dispersants should fall generally in the range of about 0.005 to 15 kg/metric ton (about 0.01 to 30 lbs/short ton), or preferably 0.05 to 2 kg/metric ton (0.1 to 4 lbs/short ton). With respect to the solvent, the concentration should fall in the range of about 0 to 12.5 kg/metric ton (about 0 to 25 lbs/short ton), or more preferably 0.05 to 2 kg/metric ton (0.1 to 4 lbs/short ton). As known in the art, "kg/metric ton" (or "pounds per short ton") refers to the weight of the additives in kilograms (or pounds), as compared to the weight in metric tons (or short tons) or oven dried pulp which is washed.

EP 0 177 113 B1

The additives can be supplied by various locations within the pulping system, such as any of the shower heads for the washers, the washer vats, the filtrate storage tank from where filtrate is recirculated through the washers, the digester, the deknitter, the repulper or the like. We have found that the additives are distributed throughout the washing system particularly well when they are added to the shower head of an intermediate washer in a series of washers, such as the second in a series of three.

The additives can be added individually, or can be premixed and added as a mixture. Preferably, for reasons of convenience and greater effectiveness, the additives are premixed. In a preferred embodiment, a mixture of from 10 to 60% nonionic surfactant, 10 to 60% polyelectrolyte dispersant, and 0 to 50% solvent to total 100% is utilized at a level of from about 0.05 to 25 kg/metric ton (about 0.1 to 50.0 pounds per short ton) of pulp (oven dried) to be washed. More preferably the mixture will comprise from 30 to 50% nonionic surfactant, from 20 to 40% polyelectrolyte dispersant, and from 20 to 40% polyelectrolyte dispersant, and from 20 to 40% solvent to total 100%. With respect to the concentration, more preferably, to achieve a suitable level of effectiveness at greater economy, the concentration will range from 0.25 to 2.5 kg/metric ton (about 0.5 to 5.0 pounds per short ton), with the mixture being added in the shower head of an intermediate washer in a series of washers.

The temperature of the wash water can range from about 37.7 to 100°C (about 100 to 212°F), preferably in the range of about 60 to 82.2°C (about 140 to 180°F).

While not wishing to be limited to any theory, we theorize the surprisingly beneficial results achieved by the method of this invention may result from the prevention of channeling within the mat, such that more of the wash water actually penetrates the mat and more efficiently displaces the liquor and impurities such as spent cooking chemicals and organic substances.

The invention will be further understood by reference to following examples which include the preferred embodiment.

Example I

Example I was performed at a typical sulfite pulping mill having three rotary cylinder vacuum washers in sequence. Pre-trial, trial and post-trial production runs were monitored. The "pre-trial" data are taken from the mill during standard production, over a period of twenty-one days. The trial occurred over a twenty-seven day period wherein the invention was utilized at the mill under otherwise standard conditions. Post-trial data were taken during approximately twenty-one days following the end of the trial, again during standard production without use of the invention. The pre-trial and post-trial data are in the nature of a control, for comparison with the data obtained from use of the method of the invention.

The trial procedure was as given in the following Table I.

TABLE I

<u>Day</u>	<u>Additive</u> kg/metric ton (lbs/short ton)
1	0.25 (0.5)
2	0.5 (1.0)
3	0.5 (1.0)
4	0.75 (1.5)
5	0.75 (1.5)
6	0.75 (1.5)
7	0.75 (1.5)
8	0.75 (1.5)
9	0.75 (1.5)
10	0.5 (1.0)
11	0.5 (1.0)
12	0.75 (1.5)
13	0.75 (1.5)
14	0.75 (1.5)
15	0.75 (1.5)
16	0.75 (1.5)
17	0.5 (1.0)
18	0.5 (1.0)
19	0.5 (1.0)
20	0.5 (1.0)
21	0.5 (1.0)
22	1.0 (2.0)
23	1.0 (2.0)
24	1.0 (2.0)
25	0 (0)
26	1.0 (2.0)
27	1.0 (2.0)

EP 0 177 113 B1

The additive used during the trial was in all cases provided to the #2 shower. It comprised a mixture of 40% by weight of a modified alcohol ethoxylate with a specific gravity of 0.97, and an activity of 100%, 30 wt-% of a low molecular weight polyacrylic acid in the form of a 48 to 50% aqueous solution, having a pH at 100% of 1.5—2.0, and about 30 wt-% tetrahydrofurfuryl alcohol having a molecular weight of about 102, a specific gravity at 20/20°C of about 1.0543. The temperature of the #2 shower solution ranged from 48.9—71.1°C (120°—160°F) during the trial.

Test results were as given in the following Table II.

TABLE II

		Pre-Trial	Trial	Post-Trial
10	Daily Production	349 (385)	362 (399)	341 (376)
	in metric tons	N=23	N=27	N=21
15	per day (and	S=33.2	S=47.9	S=37.93
	short tons per day)			
	Cooks (No./day)	6.1	6.8	7.3
20	Knotter Flow in	9478 (2504)	10409 (2750)	9720 (2568)
	litres per minute	N=10	N=26	N=21
	(a U.S. gallons	S=199	S=281	S=226
	per minute)	(98%)		(98%)
25	Knotter TDS	8.81	9.86	8.71
	(%)	N=6	N=26	N=21
		S=0.43	S=1.10	S=0.45
		(97%)		(99%)
30	#1 Washer	8.38	9.89	8.63
	Filtrate	N=6	N=26	N=21
	TDS (%)	S=0.39	S=1.07	S=0.46
		(99%)		(99%)
35	3 Washer Mat	0.475	0.399	0.480
	TDS (%)	N=6	N=26	N=21
		S=0.142	S=0.158	S=0.230
		(82%)		(82%)
40	#3 Washer Shower	2634 (696)	2858 (755)	2786 (736)
	Flow in litres	N=23	N=26	N=21
	per minute (and	S=8	S=18	S=39
	U.S. gallons			
	per minute)	(99%)		(99%)
45	Red Liquor	9.97%	10.27	9.86
	Solids (%)	N=23	N=27	N=21
		S=0.48	S=0.52	S=0.37
		(99%)		(99%)
50	Surge Tank Solids	37.87	37.44	35.75
	Loss in kg/metric	(75.74)	(74.88)	(71.51)
	ton (and lb/short	N=23	N=26	N=21
	ton)	S=17.08	S=11.34	S=8.85
		(82%)		(75%)
55	Overall	95.35%	96.37%	99.44%
	Efficiency	N=6	N=26	N=21
	(%)	S=1.22	S=1.59	S=3.24
		(85%)		(99%)
60	#3 Washer Mat	1.197	1.014	1.140
	Extractables	N=3	N=13	N=15
	%	S=0.17	S=0.14	S=0.16
65		(92%)		(90%)

EP 0 177 113 B1

The knotter TDS refers to the total dissolved solids in the liquid phase from the knotter, before washing of the pulp.

The #1 washer filtrate TDS refers to the total dissolved solids of the filtrate recovered from the first rotary cylinder vacuum washer in the series of three. A higher value indicates that a greater number of impurities have been washed from the pulp.

Number 3 washer mat TDS refers to the total dissolved solids of the mat itself which forms on washer #3. This is calculated by squeezing liquid from the mat, and testing that liquid for total dissolved solids. It is understood in the art that the composition of the liquid squeezings corresponds to the composition of the mat itself. A lower number indicates a more pure mat and is preferred.

Number 3 washer shower flow refers to the gallons per minute of wash water flowing from the head of shower number 3.

Red liquor solids refers to the weight percent of solids compared to the total weight of the red liquor. This value indicates the presence of impurities such as cooking chemicals, lignin and the like which the pump contains after digestion and which are removed during washing. A higher number indicates that more impurities have been removed from the pulp and will be recoverable.

Surge tank solids loss reflects the solids, whether cooking chemicals, lignin and the like, which are lost to recovery, that is, which have been carried over with the pulp and are therefore unrecovered. A lower value is preferred.

Overall efficiency is expressed as a percentage. A higher percentage for overall efficiency indicates greater washing efficiency and is more desirable. It is calculated by the following formula:

$$\text{Efficiency} = \frac{(\text{TDS \#1 vat}) - (\text{TDS \#3 mat})}{(\text{TDS \#1 vat}) - (\text{TDS \#3 shower})} \times 100$$

In the formula, "TDS" refers to "total dissolved solids". The TDS #1 vat refers to the total dissolved solids of the brown stock in the vat of the first in the series of washers. TDS #3 mat is a value obtained by analysis of the squeezings of the mat formed on the #3 washer. TDS #3 shower reflects the total dissolved solids of the wash water sprayed through the #3 shower head.

Number 3 washer mat extractables is expressed as the weight percent of impurities as compared to the total weight of the mat. This test was performed following the TAPPI method but using a mixture of toluene and alcohol for the extraction process, rather than benzene and alcohol. A lower number indicates fewer organic soluble impurities in the pulp or mat.

With respect to the data, N indicates the number of samples which were tested. The values obtained were then analyzed via Student T test to provide the value given in Table I. S represents the standard of deviation among the values obtained. The percentage given in parentheses indicates the confidence limits of the value.

Discussion of Results

Regression equations were developed using pre-trial and trial data with addition of the additive mixture as described in the example at a level of 0.75 kg/metric ton (1.5 lb/short ton). All work was done through multiple linear regression. Terms included in the equations are significant at a level of ($P \leq 0.05$). The regression equations are as follows:

Red Liquor Solids = $9.813 + 1.352 \times \# \text{ Mat TDS} + 1.190 \times \text{lb/short ton additive mixture used in Example} + 0.077 \times \#1 \text{ Vat TDS} - 0.167 \times \text{Cooks (No/day)}$. $R^2 = 0.8102$.

Surge Tank Solids Loss = $117.12 + (2.68 \times \text{Cooks (No/day)}) + (0.022 \times \text{Knotter flow}) - (1.78 \times \text{Efficiency}) + (0.120 \times \text{daily production})$. $R^2 = 0.5535$.

#3 Mat TDS = $6.552 + (0.030 \times \text{Knotter TDS}) - (0.066 \times \text{Efficiency}) - (0.041 \times \text{lb/short ton additive mixture used in Example})$. $R^2 = 0.6658$.

Red Liquor solids increased during the trial and decreased the post-trial period. The regression equation confirms a positive effect on red liquor solids equivalent to .19% per 0.5 kg/metric ton (1 pound/short ton) of the additive mixture, or approximately 0.30% at the average trial feed rate of 0.75 kg/metric ton (1.5 pounds/short ton).

Results of the #1 washer filtrate total dissolved solids values indicate a significant increase, showing markedly better washing results with use of the additive mixture. Knotter total dissolved solids also increased during the trial.

The total dissolved solids in the #3 washer mat decreased 17% during the trial, compared to the pre-trial and post-trial average. This value indicates surprisingly effective solids removal. The regression equation indicates that the effect of the additive mixture on #3 washer mat total dissolved solids is a decrease of .0615% when the additive mixture is used at 0.75 kg/metric ton (1.5 pound/short ton).

Overall efficiency for the three washers was found to be significantly increased during the trial. The post trial value is considered to be more reliable than the pre-trial value due to the small sample size for the pre-trial data. Efficiency was calculated from oven dried total dissolved solids.

Number 3 washer mat extractables showed a 12% decrease during the trial versus post-trial period, an unexpectedly high decrease particularly in view of the high production rates during the trial.

EP 0 177 113 B1

Example II

Example II was performed as was Example I, in the same sulfite mill using the same standard procedures. The same additive was also used.

The first pre-trial lasted 8 consecutive days, followed later by an additional 12 day pre-trial period, where data were collected regarding the standard mill operation. The trial period immediately followed the 12 day pre-trial, and lasted 5 days, while the post-trial period immediately followed the trial and lasted 3 days.

The trial procedure for Example II was as follows:

10

Day	Time	Action
1	10:30 a.m.	Sample run
15	10:45 a.m.	Additive in at 0.25 kg/metric ton (0.5 lb/short ton) to #2 shower
	1:15 p.m.	Sample run
	2:05 p.m.	Additive increased to 0.5 kg/metric ton (1.0 lb/short ton)
20	3:00 p.m.	Sample run
2	8:00 a.m.	Sample run
25	10:00 a.m.	Sample run
	10:15 a.m.	Additive increased to 0.75 kg/metric ton (1.5 lb/short ton)
	1:00 p.m.	Sample run
30	3	8:30 a.m. Sample run
	10:30 a.m.	Sample run
35	10:35 a.m.	Additive increased to 1.0 kg/metric ton (2.0 lb/short ton)
	12:30 p.m.	Sample run
40	4	8:30 a.m. Sample run
	10:30 a.m.	Sample run
	12:30 p.m.	Sample run
45	5	8:30 a.m. Sample run
	9:00 a.m.	Additive off

50 Squeezings were taken from mat #3 at various times during trial and post trial as indicated at "sample run", and were in some cases analyzed for water soluble inorganics using the TAPPI testing procedure, except that toluene and alcohol were used instead of benzene and alcohol. The results of the #3 mat squeezings analysis, in parts per million, is as follows:

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EP 0 177 113 B1

#3 Mat Squeezings

	Day: Time:	Trial			Post-Trial
		Day 1 Afternoon	Day 2 Morning	Day 4 Morning	Day 5 of Post Trial
5	Chloride	9.3	5.3	4.4	11.6
10	Sulfate	125	438	398	1350
	Aluminum	4	4	4	1.0
	Barium	2	2	2	0.7
15	Iron	3.6	7.0	3.0	6.0
	Silica as SiO ₂	17	17	17	16
20	Calcium	10	15	20	93
	Magnesium	83	200	240	1200
	Sulfite by iodine titration	40	10	40	620

The average of the #3 mat dissolved solids, expressed as a percentage by weight solids/weight of the squeezings as a whole, was as follows:

	First Pre-trial	Average of Second Pre-Trial and Post Trial	Trial
30	0.48	0.49	0.42

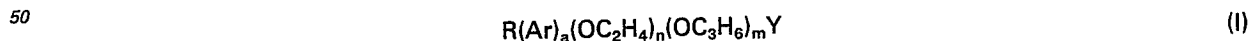
Results

Dissolved solids in the #3 mat were decreased from an average of 0.49% to 0.42% during the trial, indicating a reduction of soluble inorganics and organics. Water analysis of the #3 mat confirmed the significant reduction of inorganics during the trial. These results show a trend indicating an unexpectedly high reduction in inorganic and organic solids carried in the #3 mat during the use of the additive mixture in the #2 shower.

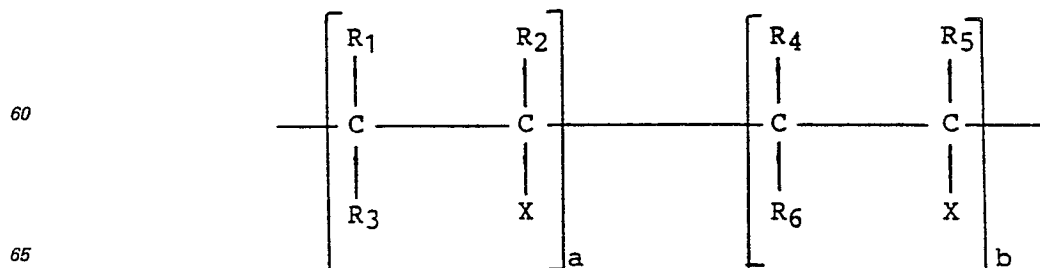
The foregoing description and Examples are exemplary of the invention.

Claims

1. An improved method of removing spent or excess cooking chemical compounds or organic contaminants from chemically or mechanically prepared virgin pulp, characterised in that it comprises (a) forming a fiber mat of chemically or mechanically prepared virgin pulp; and (b) forcing an aqueous solution comprising a substituted oxyethylene glycol non-ionic surfactant, a water soluble low molecular weight polyelectrolyte dispersant and a solvent into and through the fiber mat, wherein the non-ionic surfactant has the formula:



wherein Ar is an aromatic residue, a is zero or 1, R is an aliphatic group that contains from 6 to 24 carbon atoms when a is zero and from 6 to 18 carbon atoms when a is 1, n is from 3 to 50, m is from zero to 50, and Y is hydroxy or benzyl ether with the proviso that, when m is zero, Y is benzyl ether, and wherein the polyelectrolyte dispersant has the formula:



EP 0 177 113 B1

wherein R_1 , R_2 , R_4 and R_5 are independent and are selected from hydrogen, C_1 — C_4 lower alkyl, carboxy-substituted alkyl or mixtures thereof, R_3 and R_6 are independent and selected from hydrogen, carboxy, carboxy-substituted alkyl or mixtures thereof, X is selected from carboxy, salts and derivatives of carboxy, acetyl, hydrocarbon moieties commonly attached to free radical monomers, COOZ where Z is a monovalent metal ion or ammonium, or mixtures thereof, and the total of $a + b$ falls in the range of 15 to 1,000.

2. A method according to claim 1 wherein the forcing of the aqueous solution into and through the fiber mat comprises applying the aqueous solution to the mat and drawing the solution into and through the mat with a vacuum force.

3. A method according to claim 2 wherein the applying of the aqueous solution to the mat comprises spraying the aqueous solution onto a surface of the mat.

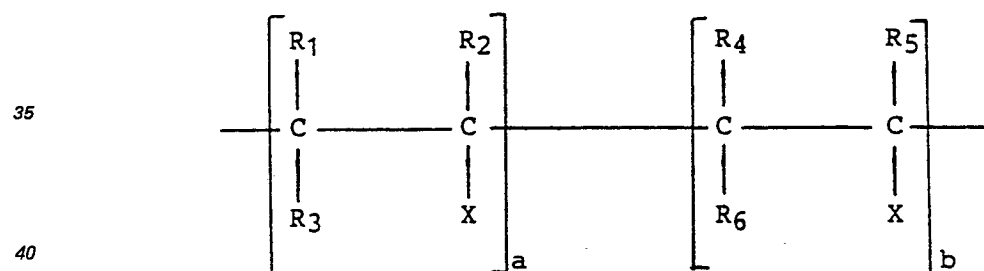
4. A method according to any of claims 1 to 3 wherein the fiber mat is formed using rotary drum vacuum washer.

5. A method according to any of claims 1 to 4 wherein the temperature of the aqueous solution is from 37.7 to 100°C (100 to 212°F).

6. A method of recovering excess or spent cooking chemical compounds or organic contaminants from an aqueous pulping medium comprising chemically or mechanically prepared virgin pulp and excess or spent chemical compounds or organic contaminants, characterised in that it comprises (a) combining the aqueous pulping medium with a substituted oxyethylene glycol non-ionic surfactant, a water soluble low molecular weight polyelectrolyte dispersant, and solvent; (b) substantially separating the chemically or mechanically prepared virgin pulp from the aqueous pulping medium; and (c) after step (b), removing excess or spent cooking chemical compounds or organic contaminants from the aqueous pulping medium; wherein the non-ionic surfactant has the formula:



wherein Ar is an aromatic residue, a is zero or 1, R is an aliphatic group that contains from 6 to 24 carbon atoms when a is zero and from 6 to 18 carbon atoms when a is 1, n is from 3 to 50, m is from zero to 50, and Y is hydroxy or benzyl ether with the proviso that, when m is zero, Y is benzyl ether, and wherein the polyelectrolyte dispersant has the formula:



wherein R_1 , R_2 , R_4 and R_5 are independent and are selected from hydrogen, C_1 — C_4 lower alkyl, carboxy-substituted alkyl or mixtures thereof, R_3 and R_6 are independent and selected from hydrogen, carboxy, carboxy-substituted alkyl or mixtures thereof, X is selected from carboxy, salts and derivatives of carboxy, acetyl, hydrocarbon moieties commonly attached to free radical monomers, COOZ where Z is a monovalent metal ion or ammonium, or mixtures thereof, and the total of $a + b$ falls in the range of 15 to 1,000.

7. A method according to claim 6 wherein in substantially separating the chemically or mechanically prepared virgin pulp from the aqueous pulping medium, the chemically or mechanically prepared virgin pulp is washed in a washing step.

8. A method according to claim 6 or 7 wherein the chemically or mechanically prepared virgin pulp is substantially separated from the aqueous pulping medium and washed by a rotary vacuum cylinder washer.

9. A method according to any of claims 1 to 8, wherein the solvent is an ethoxylated solvent.

10. A method according to claim 9 wherein the solvent comprises tetrahydrofurfuryl alcohol, ethoxylated derivative or mixtures thereof.

11. A method according to any of claims 1 to 10 wherein the substituted oxyethylene glycol non-ionic surfactant, the water soluble low molecular weight polyelectrolyte dispersant, and the solvent together are provided at a total concentration of from 0.05 to 25 kg/metric ton (0.1 to 50.0 pounds per short ton) of oven dried pulp, the total concentration being composed of 10 to 60% by weight surfactant, 10 to 60% by weight dispersant, and 20 to 50% by weight solvent.

12. A method according to claim 11 wherein the total concentration is from 0.25 to 2.5 kg/metric ton (0.5 to 5.0 pounds per short ton) of oven dried pulp.

13. A method according to any of claims 1 to 12 wherein the polyelectrolyte dispersant comprises a copolymer of maleic acid and vinyl acetate.

14. A method according to any of claims 1 to 12 wherein the polyelectrolyte dispersant comprises a polyacrylate compound.

15. A method according to claim 14 wherein the polyacrylate compound has a molecular weight in the range of 500 to 25,000.

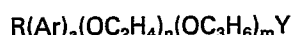
5 16. A method according to any of claims 1 to 12 wherein, the formula for the polyelectrolyte dispersant, R_1 , R_3 , R_4 , R_6 are hydrogen, R_2 and R_5 are hydrogen or methyl, and X is carboxy.

Patentansprüche

10 1. Verbessertes Verfahren zur Entfernung verbrauchter oder überschüssiger Kochchemikalien oder organischer Verunreinigungen aus chemisch oder mechanisch hergestelltem Frischzellstoffbrei, dadurch gekennzeichnet, daß es

(a) das Bilden einer Fasermatte aus chemisch oder mechanisch hergestelltem Frischzellstoffbrei und

15 (b) das zwangsweise Hinein- und Hindurchführen einer wäßrigen Lösung, enthaltend ein nicht-ionisches substituiertes Oxyethylenglycol-Tensid, ein wasserlösliches niedermolekulares Polyelektrolyt-Dispergiermittel und ein Lösungsmittel, in die Fasermatte hinein und durch diese hindurch umfaßt, wobei das nicht-ionische Tensid die Formel



20 besitzt, in der

Ar ein aromatischer Rest ist,

a null oder 1 ist,

R eine aliphatische Gruppe ist, die 6 bis 24 Kohlenstoff-Atome enthält, wenn a null ist, und 6 bis 18

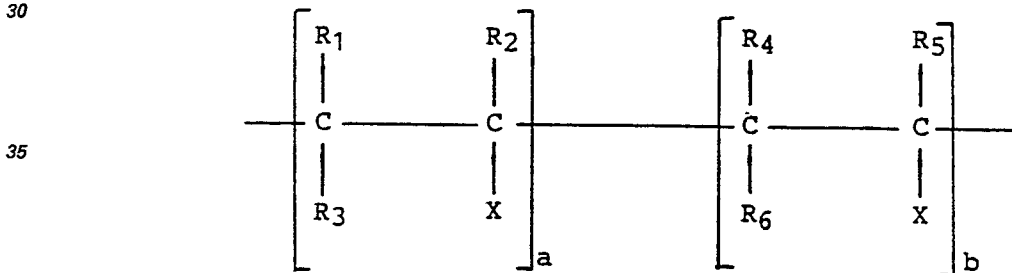
25 Kohlenstoff-Atome enthält, wenn a 1 ist,

n 3 bis 50 ist,

m null bis 50 ist und

Y Hydroxy oder Benzylether ist, mit der Maßgabe, daß, wenn m null ist, Y Benzylether ist, worin das Polyelektrolyt-Dispergiermittel die Formel

30



40 besitzt, in der

R_1 , R_2 , R_4 und R_5 unabhängig voneinander sind und aus Wasserstoff, C_1 — C_4 -Niederalkyl, carboxy-substituiertem Alkyl oder deren Mischungen ausgewählt sind,

R_3 und R_6 unabhängig voneinander sind und aus Wasserstoff, Carboxy, carboxy-substituiertem Alkyl oder deren Mischungen ausgewählt sind,

45 X aus Carboxy, Salzen und Derivaten von Carboxy, Acetyl, Kohlenwasserstoff-Einheiten, die gewöhnlich an freie Radikal-Monomere gebunden sind, COOZ, worin Z ein einwertiges Metall-Ion oder Ammonium ist, oder deren Mischungen ausgewählt ist und der Gesamtwert $a + b$ in den Bereich von 15 bis 1000 fällt.

50 2. Verfahren nach Anspruch 1, worin das zwangsweise Hinein- und Hindurchführen der wäßrigen Lösung in die Fasermatte hinein und durch diese hindurch das Aufbringen der wäßrigen Lösung auf die Matte und das Ziehen der Lösung in die Matte hinein und durch diese hindurch mittels einer Vakuum-Kraft umfaßt.

3. Verfahren nach Anspruch 2, worin das Aufbringen der wäßrigen Lösung auf die Matte das Sprühen der wäßrigen Lösung auf eine oberfläche der Matte umfaßt.

55 4. Verfahren nach irgendeinem der Ansprüche 1 bis 3, worin die Fasermatte mittels eines Drehtrommel-Vakuumwäschers gebildet wird.

5. Verfahren nach irgendeinem der Ansprüche 1 bis 4, worin die Temperatur der wäßrigen Lösung 37,7°C bis 100°C (100°F bis 212°F) beträgt.

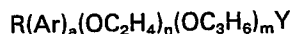
60 6. Verfahren zur Rückgewinnung überschüssiger ode verbrauchter Kochchemikalien oder organischer Verunreinigungen aus einem wäßrigen Zellstoffmedium, das chemisch oder mechanisch hergestellten Frischzellstoffbrei und überschüssige oder verbrauchte chemische Verbindungen oder organische Verunreinigungen enthält, dadurch gekennzeichnet, daß es

65 (a) das Vereinigen des wäßrigen Zellstoff-Mediums mit einem nicht-ionischen substituierten Oxyethylenglycol-Tensid, einem wasserlöslichen, niedermolekularen Polyelektrolyt-Dispergiermittel und einem Lösungsmittel,

EP 0 177 113 B1

(b) im wesentlichen das Abtrennen des chemisch oder mechanisch hergestelltem Frischzellstoffbreis aus dem wäßrigen Zellstoffmedium und

(c) nach dem Schritt (b), das Entfernen der überschüssigen oder verbrauchten Kochchemikalien oder organischen Verunreinigungen aus dem wäßrigen Aufschlußmedium
5 umfaßt, wobei das nicht-ionische Tensid die Formel



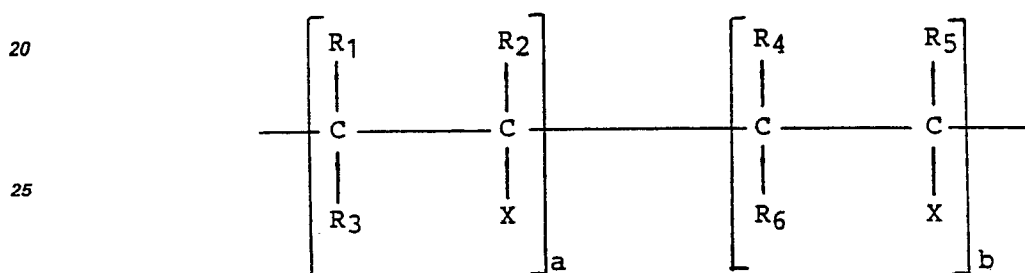
besitzt, in der

10 Ar ein aromatischer Rest ist,
a null oder 1 ist,

R eine aliphatische Gruppe ist, die 6 bis 24 Kohlenstoff-Atome enthält, wenn a null ist, und 6 bis 18 Kohlenstoff-Atome enthält, wenn a 1 ist,
n 3 bis 50 ist,

15 m null bis 50 ist und

Y Hydroxy oder Benzylether ist, mit der Maßgabe, daß, wenn m null ist, Y Benzylether ist, worin das Polyelektrolyt-Dispergiermittel die Formel



30 besitzt, in der

R₁, R₂, R₄ und R₅ unabhängig voneinander sind und aus Wasserstoff, C₁—C₄-Niederalkyl, carboxy-substituiertem Alkyl oder deren Mischungen ausgewählt sind,

R₃ und R₆ unabhängig voneinander sind und aus Wasserstoff, Carboxy, carboxy-substituiertem Alkyl oder deren Mischungen ausgewählt sind,

35 X aus Carboxy, Salzen und Derivaten von Carboxy, Acetyl, Kohlenwasserstoff-Einheiten, die gewöhnlich an freie Radikal-Monomere gebunden sind, COOZ, worin Z ein einwertiges Metall-Ion oder Ammonium ist, oder deren Mischungen ausgewählt ist und der Gesamtwert a + b in den Bereich von 15 bis 1000 fällt.

7. Verfahren nach Anspruch 6, worin bei dem wesentlichen Abtrennen des chemisch oder mechanisch hergestellten Frischzellstoffbreis aus dem wäßrigen Zellstoffmedium der chemisch oder mechanisch hergestellte Frischzellstoffbrei in einem Waschschrift gewaschen wird.

8. Verfahren nach Anspruch 6 oder Anspruch 7, worin der chemisch oder mechanisch hergestellte Frischzellstoffbrei im wesentlichen von dem wäßrigen Zellstoffmedium abgetrennt und mittels eines Drehzylinder-Vakuumwäschers gewaschen wird.

45 9. Verfahren nach irgendeinem der Ansprüche 1 bis 8, worin das Lösungsmittel ein ethoxyliertes Lösungsmittel ist.

10. Verfahren nach Anspruch 9, worin das Lösungsmittel Tetrahydrofurfurylalkohol, ein ethoxyliertes Derivat oder deren Gemische umfaßt.

50 11. Verfahren nach irgendeinem der Ansprüche 1 bis 10, worin das nicht-ionische substituierte Oxyethylenglycol-Tensid, das wasserlösliche, niedermolekulare Polyelektrolyt-Dispergiermittel und das Lösungsmittel zusammen in einer Gesamt-Konzentration von 0,05 bis 25 kg/metrisch Tonne (0,1 bis 50,0 pounds pro short ton) ofengetrockneter Zellstoffmasse bereitgestellt werden, wobei die Gesamt-Konzentration aus 10 bis 60 Gew.-% Tensid, 10 bis 60 Gew.-% Dispergiermittel und 20 bis 50 Gew.-% Lösungsmittel zusammengesetzt ist.

55 12. Verfahren nach Anspruch 11, worin die Gesamt-Konzentration von 0,25 bis 2,5 kg/metrische Tonne (0,5 bis 5,0 pounds pro short ton) ofengetrockneter Zellstoffmasse beträgt.

13. Verfahren nach irgendeinem der Ansprüche 1 bis 12, worin das Polyelektrolyt-Dispergiermittel ein Copolymer aus Maleinsäure und Vinylacetat umfaßt.

60 14. Verfahren nach irgendeinem der Ansprüche 1 bis 12, worin das Polyelektrolyt-Dispergiermittel eine Polyacrylat-Verbindung umfaßt.

15. Verfahren nach Anspruch 14, worin die Polyacrylat-Verbindung ein Molekulargewicht im Bereich von 500 bis 25 000 besitzt.

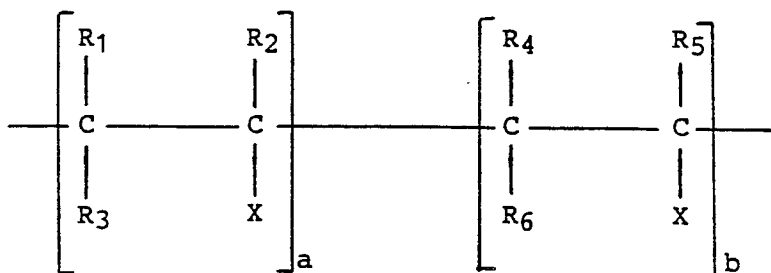
16. Verfahren nach irgendeinem der Ansprüche 1 bis 12, worin in der Formel für Polyelektrolyt-Dispergiermittel R₁, R₃, R₄ und R₆ Wasserstoff sind, R₂ und R₅ Wasserstoff oder Methyl sind und X Carboxy ist.

Revendications

1. Un procédé perfectionné pour éliminer les composés chimiques de cuisson usés ou en excès ou les contaminants organiques d'une pâte vierge préparée chimiquement ou mécaniquement, caractérisé en ce qu'il comprend (a) la formation d'une nappe de fibres de pâte vierge préparée chimiquement ou mécaniquement; et (b) le passage à force d'une solution aqueuse, comprenant un agent tensio-actif non ionique de type oxyéthylèneglycol substitué, un polyélectrolyte dispersant de bas poids moléculaire soluble dans l'eau et un solvant, dans et à travers la nappe de fibres, l'agent tensio-actif non ionique répondant à la formule:



dans laquelle Ar est un résidu aromatique, a est zéro ou 1, R est un groupe aliphatique qui contient de 6 à 24 atomes de carbone lorsque a est zéro et de 6 à 18 atomes de carbone lorsque a est 1, n vaut de 3 à 50, m vaut de zéro à 50 et Y est un hydroxy ou un benzyléther, sous réserve que lorsque m est zéro, Y est un benzyléther, et où le polyélectrolyte dispersant a pour formule:



dans laquelle R_1 , R_2 , R_4 et R_5 sont indépendants et sont choisis parmi un hydrogène, un alcoyle inférieur en C_1-C_4 , un alcoyle carboxy-substitué ou leurs mélanges, R_3 et R_6 sont indépendants et choisis parmi un hydrogène, un carboxy, un alcoyle carboxy-substitué ou leurs mélanges, X est choisi parmi un carboxy, les sels et dérivés d'un carboxy, un acétyl, des fragments hydrocarbures fixés ensemble à des monomères radicalaires, COOZ où Z est un ion de métal monovalent ou un ammonium, ou leurs mélanges, et le total de $a + b$ se situe dans la gamme de 15 à 1 000.

2. Un procédé selon la revendication 1, dans lequel le passage à force de la solution aqueuse dans et à travers la nappe de fibres comprend l'application de la solution aqueuse à la nappe et l'attraction de la solution dans et à travers la nappe avec une force créée par le vide.

3. Un procédé selon la revendication 2, dans lequel l'application de la solution aqueuse à la nappe comprend la pulvérisation de la solution aqueuse sur une surface de la nappe.

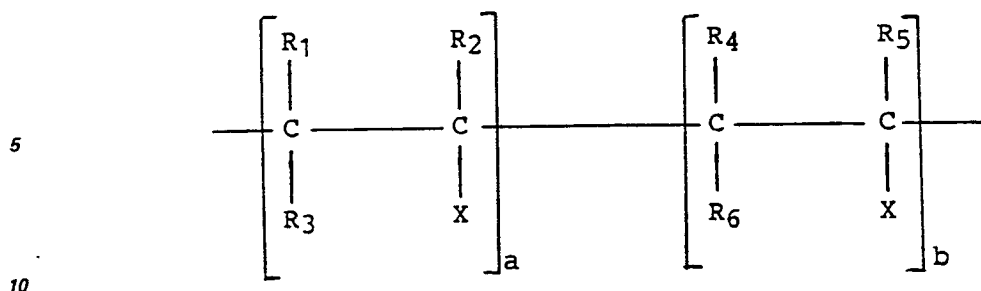
4. Un procédé selon l'une quelconque des revendications 1 à 3, dans lequel la nappe de fibres est formée à l'aide d'une pile laveuse aspirante à tambour rotatif.

5. Un procédé selon l'une quelconque des revendications 1 à 4, dans lequel la température de la solution aqueuse est de 37,7 à 100°C (100 à 212°F).

6. Un procédé de récupération des composés chimiques de cuisson en excès ou usés ou des contaminants organiques d'un milieu aqueux de fabrication de la pâte, comprenant de la pâte vierge préparée chimiquement ou mécaniquement et des composés chimiques en excès ou usés, ou des contaminants organiques, caractérisé en ce qu'il comprend (a) la combinaison du milieu aqueux de formation de la pâte avec un agent tensio-actif non ionique de type oxyéthylèneglycol substitué, un polyélectrolyte dispersant de bas poids moléculaire soluble dans l'eau et un solvant; (b) une séparation importante de la pâte vierge préparée chimiquement ou mécaniquement d'avec le milieu aqueux de formation de la pâte; et (c) après le stade (b), l'élimination des composés chimiques de cuisson en excès ou usés ou des contaminants organiques du milieu aqueux de formation de la pâte; dans lequel l'agent tensio-actif non ionique répond à la formule:



dans laquelle Ar est un résidu aromatique, a est zéro ou 1, R est un groupe aliphatique qui contient de 6 à 24 atomes de carbone lorsque a est zéro et de 6 à 18 atomes de carbone lorsque a est 1, n vaut de 3 à 50, m vaut de zéro à 50 et Y est un hydroxy ou un benzyléther, sous réserve que lorsque m est zéro, Y est un benzyléther, et où le polyélectrolyte dispersant a pour formule:



dans laquelle R_1 , R_2 , R_4 et R_5 sont indépendants et sont choisis parmi un hydrogène, un alcoyle inférieur en C_1-C_4 , un alcoyle carboxy-substitué ou leurs mélanges, R_3 et R_6 sont indépendants et choisis parmi un hydrogène, un carboxy, un alcoyle carboxy-substitué ou leurs mélanges, X est choisi parmi un carboxy, les sels et dérivés d'un carboxy, un acétyl, des fragments hydrocarbures fixés ensemble à des monomères radicalaires, $COOZ$ où Z est un ion de métal monovalent ou un ammonium, ou leurs mélanges, et le total de $a + b$ se situe dans la gamme de 15 à 1 000.

7. Un procédé selon la revendication 6, dans lequel, dans la séparation importante de la pâte vierge préparée chimiquement ou mécaniquement d'avec le milieu aqueux de formation de la pâte, la pâte vierge préparée chimiquement ou mécaniquement est lavée dans un stade de lavage.

8. Un procédé selon la revendication 6 ou 7, dans lequel la pâte vierge préparée chimiquement ou mécaniquement est essentiellement séparée d'avec le milieu aqueux de formation de la pâte à l'aide d'une pile laveuse aspirante à cylindre rotatif.

9. Un procédé selon l'une quelconque des revendications 1 à 8, dans lequel le solvant est un solvant éthoxylé.

10. Un procédé selon la revendication 9, dans lequel le solvant comprend de l'alcool tétrahydrofur, un dérivé éthoxylé ou leurs mélanges.

11. Un procédé selon l'une quelconque des revendications 1 à 10, dans lequel l'agent tensio-actif non ionique de type oxyéthylèneglycol substitué, le polyélectrolyte dispersant de bas poids moléculaire soluble dans l'eau et le solvant réunis ont une concentration totale de 0,05 à 25 kg/t (0,1 à 5,0 lb/short ton) de pâte séchée à l'étuve, la concentration totale étant constituée de 10 à 60% en poids d'agent tensio-actif, 10 à 60% en poids de dispersant et 20 à 50% en poids de solvant.

12. Un procédé selon la revendication 11, dans lequel la concentration totale est de 0,25 à 2,5 kg/t (0,5 à 5,0 lb/short ton) de pâte séchée à l'étuve.

13. Un procédé selon l'une quelconque des revendications 1 à 12, dans lequel le polyélectrolyte dispersant comprend un copolymère d'acide maléique et d'acétate de vinyle.

14. Un procédé selon l'une quelconque des revendications 1 à 12, dans lequel le polyélectrolyte dispersant comprend un composé polyacrylate.

15. Un procédé selon la revendication 14, dans lequel le composé polyacrylate a un poids moléculaire dans la gamme de 500 à 25 000.

16. Un procédé selon l'une quelconque des revendications 1 à 12, dans lequel, dans la formule du polyélectrolyte dispersant, R_1 , R_3 , R_4 et R_6 sont un hydrogène, R_2 et R_5 sont un hydrogène ou un méthyle et X est un carboxy.