

METHOD AND APPARATUS FOR FORMING A SUSPENSION OF PARTLY DISSOLVED FLY ASH IN A MINERAL ACID

The present invention relates to a method and apparatus for forming a suspension of partly dissolved fly ash in a mineral acid.

5 **Background**

Fly ash is a fine powdered waste product entrained in the combustion gases from combustion and incineration processes. The fly ash contains a mixture of mineral oxides and a range of other chemical compounds which may have a detrimental effect on the environment. Present environmental standards therefore generally
10 forbid discharging the fly ash to the atmosphere.

Thus, present day combustion/incineration plants typically apply capturing technology to separate the fly ash out of the combustion gases before venting the gases to the atmosphere. This produces huge amounts of captured fine powdered ash which must be disposed of. For example, NOAH AS, an industrial waste disposal
15 company operating in Scandinavia receives more than 300 000 metric tonnes of fly ash every year from incineration plants.

Due to the content of heavy metals, fly ash from incinerators is considered a harmful waste and is subject to environmental restrictions requiring specially secured waste disposals for deposition of the fly ash. For example, fly ash from a
20 municipal incineration plant in Oslo, Norway, is treated with sulphuric acid to encapsulate the fly ash in gypsum, which then is sent to disposal at a landfill for hazardous waste. However, the disposal of huge amounts of fly ash in landfill deposits for harmful waste is problematic both from an economical and a resource point of view.

Fly ash from incinerators of municipal waste contains relatively large concentrations of volatile metals, such as e.g. zinc vaporised in the incineration and condensed to salts, oxides, silicates or other compounds in the fly ash. Fly ash may contain around 1.5 wt% zinc such that disposal of fly ash in landfills often buries
25 vast amounts of zinc. This is problematic from a resource point of view because it is estimated that known global reserves on zinc containing ores holds around 220
30 million metric tonnes of zinc, while the annual global zinc consumption is about 13 million metric tonnes, of which about 1/3 is recycled. These figures inform that it is necessary to significantly increase the recycle ratio to secure future access to zinc.

To extract and recycle the volatile metals in the fly ash also makes sense from an
35 economical point of view, because the deposition of vast amounts fly ash per year in leakage secured landfills is costly. Without the volatile metals, the fly ash may no longer be classified as environmental hazardous waste and may be disposed in ordinary land-fills or applied as raw material for other products, for example foam

glass aggregate. In addition, the extracted metal products can be developed to commercial commodities enabling generating revenue from waste fractions.

Prior art

5 From US 5 512 257 it is known a process where fly ash from incineration plants is treated with an acidic aqueous solution for the purpose of removing heavy metals. The residual solid matter is filtered. The filtration residue is treated with a precipitating agent for heavy metals. The thereby obtained inert residue can be directly dumped.

10 US 4 617 180 discloses a process for purifying flue gases developed during the combustion of waste materials and containing solids of various compositions as well as pollutant gases and heavy metals in gaseous form, wherein the primarily solid components are removed from the flue gas in a separating system and are collected as flue ashes and wherein the flue gas is washed for the purpose of binding the pollutant gases of the residual solids components, particularly in the slag. It is the object of the present invention to provide a process wherein, during
15 flue gas purification, the gaseous heavy metal contaminants as well as the heavy metal contaminants which are present in the flue ash and in the slag in soluble form, can be removed from the flue gas purification process in the most compact form, with simultaneously achieving good, or even improved, initial purification of HCl, SO₂, and NO_x. All other residues should be recyclable for further use. The
20 remainder of heavy metal components which are not bound in the slag are isolated from the slag by a subsequent weakly acid treatment.

From JP 2001/113242 it is known a method including a salt elution process in which fly ash is added with a mineral acid to be converted into slurry, a heavy metal
25 separation process in which the slurry is neutralized to be separated into precipitates containing heavy metals and filtrate containing salts, a dissolved oxygen removing process for removing dissolved oxygen from the filtrate, an iron salt coprecipitation process in which an iron (II) salt is added to adjust pH at 8-11, preferably at 8-9 and 9-11 in turn in two stages, to coprecipitate selenium and other heavy metals with
30 iron hydroxide, a precipitate washing process for washing the heavy metal-containing precipitates, and a heavy metal recovery process in which the washed precipitates, after being repulped with a mineral acid, is neutralized again for solid-liquid separation. The filtrate in the heavy metal recovery process is used as water to make the slurry in the salt elution process.

35 JP 2003/181413 discloses a treatment method of fly ash comprising a heavy metal elution process in which preferably a solution of sodium hydroxide as an alkali agent is added to the fly ash and, thereby, the fly ash is slurried, a calcium separating process in which the slurry is diluted and is fractionated into a calcium-containing deposit and a heavy metal-containing filtrate, a heavy metal separating

process in which a mineral acid is added to the heavy metal- containing filtrate, thereby, the filtrate is adjusted to pH 7-10 and, thereafter, the heavy metal is fractionated to hydroxides or sulfates or a lead separating process in which sulfuric acid as the mineral acid is added to the heavy metal- containing filtrate and the filtrate is adjusted to $\leq$ pH 5, preferably $\leq$ pH 4 and, thereafter, is fractionated to a lead-containing deposit and a zinc-containing filtrate and a zinc separating process in which the alkali agent is added to the zinc-containing filtrate, the zinc-containing filtrate is adjusted to pH 7-10, preferably, to pH 8-9 and, thereafter, is fractionated to a zinc-containing deposit and a salt-containing filtrate.

From JP 2005/272955 it is known a treatment method for fly ash where heavy metals can be recovered from fly ash generated at the time of incinerating waste such as shredder dust at a high recovery rate, and can be recycled for a refining raw material or the like. After an acid leaching stage where acid such as hydrochloric acid is added to slurry comprising fly ash showing acidity when being dissolved in pure water such as fly ash generated at the time of incinerating waste such as shredder dust to regulate its pH to $\leq$ 3, preferably to 1 to 2, thus heavy metals are ionized, if required, a sulfurizing stage where a sulfurizing agent is added thereto is performed, and a heavy metal recovering stage where heavy metals such as copper and zinc are separated and recovered by floatation is performed.

JP 2007/186761 discloses a valuable metal recovery method comprises the steps of: dechlorination-washing fly ash containing the valuable metal and chlorine with a washing liquid with a pH of 10 to 12; mixing the dechlorinated fly ash, a zinc-containing raw material, a flux, and coal; drying the mixture; crushing the dried product; forming the crushed product into an agglomerated ore; and melt-reducing the agglomerated ore. Alternatively, the valuable metal recovery method comprises the steps of: recovering zinc as crude zinc oxide by melt-reduction of the agglomerated ore; further dechlorination-washing the recovered crude zinc oxide with a washing liquid with a pH of 10 to 12; and using the dechlorinated crude zinc oxide in zinc refining.

Objective of the invention

The main objective of the invention is to provide a method and reactor for forming a suspension of partly dissolved fly ash in a mineral acid.

A further objective of the invention is to provide a method and reactor which utilises empirical information of how the pH and temperature of the mineral acid responds to added fly ash to regulate the addition of fly ash to an optimized dissolution degree of the one or metals which are to be extracted from the fly ash.

A further objective of the invention is to provide a method and plant for extracting one or more metals from fly ash utilising the method according to the invention for forming a suspension of partly dissolved fly ash in a mineral acid.

5 Description of the invention

The invention may be considered as a method for automatic regulation of the addition of solid fly ash to a mineral acid to achieve an optimized dissolution degree of the fly ash, and a reactor for executing the automatic regulation and obtain a slurry with an optimized content of dissolved metals to be extracted in downstream processing.

Thus, in a first aspect, the invention relates to a method for forming a suspension of partly dissolved fly ash in an aqueous mineral acid solution, wherein the method comprises the following process steps:

- 1) supplying a mass, L , of an aqueous mineral acid solution to a dissolution reactor and then measuring and registering the initial temperature, T_0 , and initial, pH_0 , of the aqueous mineral acid solution,
- 2) set $n = 1$, where n is an integer, and select a target pH, pH_{targ} , to be reached in the suspension,
- 3) for $n \leq N$, where N is an integer from 3 or higher, execute steps i), ii) and iii) in successive order:
 - i) adding and mixing an amount S_n of solid fly ash into the aqueous mineral acid solution,
 - ii) when the amount S_n is dissolved, measuring and registering the temperature and pH of the aqueous mineral acid solution at this n 'th step as T_n and pH_n , respectively, and determining and registering an accumulated amount of fly ash, $S_{n, \text{acc}}$, being added to the aqueous mineral acid solution at this n 'th step, and
 - iii) set $n = n+1$ and go to step i),
- 4) apply the registered temperatures T_0 to T_n and the registered pH-values pH_0 to pH_n , and the registered amounts of added fly ash, $S_{n, \text{acc}}$, and equations:

$$\Gamma_n^2 = \beta_1 - \beta_2 \Delta T_n \quad \text{and} \quad \Delta T_n = \gamma_1 \left(\frac{S_{n, \text{acc}}}{L} \right) - \gamma_2 \left(\frac{S_{n, \text{acc}}}{L} \right)^2,$$

where $\Gamma_n = 10^{-pH_n}$, $\Delta T_n = T_n - T_0$, and β_1 , β_2 , γ_1 , and γ_2 are parameters, to determine the parameters β_1 , β_2 , γ_1 , and γ_2 by regression analysis,

- 5) determining a total amount, S_{tot} , of solid fly ash to be added to the aqueous mineral acid solution from the relation:

$$S_{tot} = \frac{L}{2\gamma_2} [\gamma_1 + \sqrt{\gamma_1^2 + 4\gamma_2 \left(\frac{\beta_1 - (10^{-pH_{targ}})^2}{\beta_2} \right)}],$$

and

6) if $S_{n, acc} < S_{tot}$, execute steps j) and jjj):

- 5 j) adding and mixing an amount S_n into the aqueous mineral acid solution,
- jjj) if $S_{n, acc} < S_{tot}$, set $n = n+1$ and go to step j),
- or else:
- jjjj) stop feeding and mixing solid fly ash into the aqueous mineral acid solution.
- 10 Fly ash is a mixture of particulate matter captured in flue gases exiting a combustion/incineration process, i.e. particles being sufficiently small and light to be entrained and carried away with the flue gas. Fly ash consist thus of a mixture of particulate condensed matter, particles of ash, and particles of more or less unburnt remains of the fuel being combusted. The composition of the fly ash is thus highly
- 15 dependent on the process parameters of the combustion/incineration process and the fuel being combusted. The invention according to the first aspect of the invention is not tied to any specific source of fly ash, it may apply fly ash formed in any combustion process. However, an especially suited and preferred fly ash is fly ash from incineration of municipal waste (hereby denoted; MSWI fly ash) due to its
- 20 relatively high content of heavy metals and other metals which may be converted to valuable products/commercial commodities, and further because MSWI-fly ash typically also contains relatively high fractions of chlorides which may be converted to commercial commodities such as road salt or other products. MSWI-fly ash is particularly varying in composition due to relatively huge variations in the
- 25 composition of the waste being incinerated, however, the following elements are usually present in various amounts, ref [1]; Al, Ca, Cl, K, Na, Si, and O, and heavy metals such as; Cr, Cu, Pb, and Zn.

30 Fly ash contains both soluble compounds, typically salts such as oxides and chlorides of the above-mentioned elements (and other elements), and non-soluble compounds. The solubility of several typical salts in the fly ash is relatively low in neutral water (pH around 7). For example, MSWI-fly ash added to liquid water in a solid to liquid weight ratio of 1 : 10, only dissolves a minor fraction of the salt content and the pH stabilises around pH = 12, depending on the CaO content in the fly ash. However, at lower pH values, the solubility of these salts increases significantly.

35 The stronger the acid, the higher dissolution rates of the salts. Thus, the term “forming a suspension of partly dissolved fly ash” as used herein, means that an amount of fly ash is added to an aqueous mineral acid solution having a pH of 2 or less (before addition of the fly ash) to enable a dissolution rate of the salt content of the fly ash. The mineral acid may be any mineral acid, such as e.g. hydrochloric

acid, nitric acid, nitrous acid, sulphuric acid, sulphurous acid, etc. For example, MSWI-fly ash added to an 1 M aqueous hydrochloric acid in a solid to liquid weight ratio of 1 : 10, will result in substantial dissolution of the salt content of the fly ash and the pH becomes stabilized in the range of from pH = 1 to pH = 3, depending on the composition of the fly ash.

The major part of the soluble salts in fly ash is basic and consumes mineral acid when dissolving. The pH of the suspension increases in proportion to how much base being dissolved. However, the solubility of at least some of the basic compounds in the fly ash decreases significantly when the pH becomes higher.

Thus, the amount of fly ash being added to the aqueous mineral acid solution may advantageously be balanced between, on one hand, the need for adding as much fly ash as possible to obtain as high concentration as possible in the suspension of the compounds intended to be extracted from the fly ash, and on the other hand, to restrict the amount of added fly ash to give an over-stoichiometric amount of acid to ensure that the suspension is sufficiently acidic to dissolve most or all of the dissolvable compounds in the fly ash. I.e. to avoid the pH of the suspension to become too high and cause insufficient dissolution of added fly ash. The optimal balance is obtained when adding as much fly as possible ash to the aqueous mineral acid solution, but not more than which is effectively dissolved. The term “target pH, pH_{targ}” as used herein represents the pH-value of a suspension after having added an optimally balanced amount of solid fly ash to an aqueous mineral acid and the dissolvable compounds of the fly ash have reached chemical equilibrium with the mineral acid. Different salts have different solubility as function of pH. The target pH which should be selected depends therefore on which metal salt(s) in the solid fly ash is being intended to be dissolved and later extracted by precipitation.

However, in practice, target value representing the optimal balance may advantageously be chosen to be in one of the following ranges; of from pH_{targ} = 1.0 to pH_{targ} = 4.0, more preferably of from pH_{targ} = 1.5 to pH_{targ} = 3.0, more preferably of from pH_{targ} = 1.5 to pH_{targ} = 2.5, and most preferably of from pH_{targ} = 1.5 to pH_{targ} = 2.0.

The simplest thinkable method for regulating the addition of solid fly ash is to apply one or more pH-sensor(s) in the mineral acid and terminate the feed of solid fly ash when measuring a pH equal to the selected target pH, pH_{targ}. However, pH measurements in suspensions have shown to be prone to systematic error readings, especially at low pH-values.

Another simple approach for regulating the amount of solid fly ash to the mineral acid is to apply the acid balance which may be defined by the relation:

$$pH = -\log_{10} \left(C_0 - \frac{C_{BS}}{L} \right) \quad (1)$$

where C_0 is the initial concentration of mineral acid in the aqueous mineral acid solution, L is amount (by mass) of the aqueous mineral acid solution, C_B is the concentration of dissolvable base in the solid fly ash, and S is amount (by mass) of solid fly ash added to the suspension. From relation (1) the total amount of solid fly ash, S_{tot} , to be added to obtain pH_{targ} is:

$$S_{tot} = \frac{L}{C_B} [C_0 - 10^{-pH_{targ}}] \quad (2)$$

Eqn. (2) provides satisfactory estimates of total amount of solid fly ash, S_{tot} , to be added to the mineral acid to arrive at the selected target pH when the strength of the mineral acid and the composition, i.e. the contents of and which basic salts being present in the solid fly ash, are known prior to the formation of the suspension of partly dissolved fly ash in an aqueous mineral acid solution. However, the exact strength of the mineral acid and/or content of basic salts in the fly ash is not always known, especially when applying waste fractions as the raw materials. Then there may be significant variations in the composition of the fly ash being added during the formation of the suspension, i.e. the composition of addition S_n may be significantly different from the composition of addition S_{n+1} , especially when applying fly ash from municipal waste incineration.

In such cases, it is highly advantageous to be able to regulate the amount of fly ash being added without prior knowledge of acid strength and/or composition of the fly ash, and even more advantageous if the regulation of amount of added fly ash may automatically adapts itself in relation to varying (and unknown) fly ash composition during addition. Besides, also utilising the acid balance (eqn. 2) to regulate the feed also includes relying on pH measurements which may be prone to systematic errors. Thus, the method according to the first aspect of the invention utilises both how the pH and the temperature of the aqueous mineral acid solution responds to solid fly ash being added and mixed into it to regulate the feed of fly ash.

The dissolution of fly ash in a mineral acid is exothermic, the temperature of the suspension being formed increases in proportion to the content of basic salts being added together with the fly ash. The temperature increase resulting from the exothermic dissolution of compounds in the fly ash is usually not higher than around 20 – 25 °C. At such moderate temperature increases, it is a good approximation to neglect the heat loss to the environment and set that the exothermic heat of dissolution equals the increase in heat content of the suspension, such that the temperature increase, ΔT , in the suspension is given by the heat balance:

$$\Delta T = a(C_0 - \frac{C_B S}{2L}) \frac{S}{L} \quad (3)$$

where “a” is a constant to be determined, L is amount (by mass) of the aqueous mineral acid solution, C_B is the concentration of dissolvable base in the solid fly ash, and S is amount (by mass) of solid fly ash added to the suspension, and C_0 is the initial concentration of mineral acid in the aqueous mineral acid solution.

5 Relation (3) may also be given as:

$$\Delta T = a\left(\Gamma + \frac{C_B S}{2L}\right) \frac{S}{L} \quad (4)$$

where $\Gamma = C_0 - \frac{C_B S}{L} = 10^{-\text{pH}}$, to illustrate that the pH increase given by relation (1) is connected to the temperature increase given in relation (3).

10 The acid balance function, eqn. (1) is numerically challenging because at low pH relatively large changes in added fly ash only lead to very small changes in pH, whereas at higher pH, only small changes in added fly ash may lead to large changes in pH. This may be alleviated by combining the acid balance and heat balance, i.e. equations eqn. (1) and eqn. (3) can be combined to give:

$$15 \quad \text{pH} = -\frac{1}{2} \log_{10} \left(C_0^2 - \frac{2C_B \Delta T}{a} \right) \quad (5)$$

This expresses pH as a function of ΔT , not involving the added amount of S. Eqn. (5) may be parametrized and given on form:

$$\Gamma^2 = \beta_1 - \beta_2 \Delta T \quad (6)$$

20 where β_1 and β_2 are parameters to be determined. Likewise, the heat balance may be parametrized and given on form:

$$\Delta T = \gamma_1 \left(\frac{S}{L} \right) - \gamma_2 \left(\frac{S}{L} \right)^2 \quad (7)$$

25 where γ_1 and γ_2 are parameters to be determined. The parametrised eqns. (6) and (7) may be applied together with the registered measurements of the pH and temperature of the aqueous mineral acid solution and the registered amounts of solid fly ash, S, added to the aqueous mineral acid solution to determine the parameters β_1 , β_2 , γ_1 , and γ_2 by non-linear regression analysis. When these parameters are determined, eqns. (6) and (7) may be combined and reformulated to provide an expression of added amount of solid fly ash as a function of the pH of the suspension/aqueous mineral acid solution. This relation may then be applied to
30 determine the total amount, S_{tot} , to be added to the aqueous mineral acid solution to arrive at the selected target pH, pH_{targ} :

$$S_{\text{tot}} = \frac{L}{2\gamma_2} \left[\gamma_1 + \sqrt{\gamma_1^2 + 4\gamma_2 \left(\frac{\beta_1 - (10^{-\text{pH}_{\text{targ}}})^2}{\beta_2} \right)} \right] \quad (8)$$

The determination of the parameters β_1 , β_2 , γ_1 , and γ_2 by regression analysis should include at least three measurements of the pH and temperature as a function of added amount of solid fly ash to be able to obtain sufficiently accurate parameters to provide a representative determination of S_{tot} . The method according to first aspect of the invention sets thus $N = 3$ or higher to enable using the regression analysis with at least three successive additions of an amount S_n of fly ash and subsequent measurements of the corresponding pH and temperature change in the aqueous mineral acid solution before applying the regression analysis and subsequent use of eqn. (8) to determine S_{tot} . However, the regression analysis becomes refined and more reliable the more measurements being used in the analysis, such that the number N of successive additions of S_n and subsequent measurements of the corresponding pH and temperature change in the aqueous mineral acid solution may advantageously be any integer from (and including) 3 to (and including) 10, preferably from 4 to 8, more preferably from 5 to 7.

The N additions of S_n should not lead to the accumulated amount $S_{n, \text{acc}}$ becoming larger than the S_{tot} required to reach the selected target pH. The amount S_n should thus not be larger than about 1/10 of S_{tot} , preferably smaller. One way of obtaining this objective is simply to make an educated guess of total amount to be added, S_{est} , and set S_n to be k 'th fraction of S_{est} , i.e. set:

$$S_n = \frac{S_{\text{est}}}{k} \quad (9)$$

where k is an integer selected to be from (and including) 5 to (and including) 100, preferably from 7 to 75, more preferably from 10 to 50, and more preferably from 20 to 30.

Alternatively, the estimation of total amount to be added, S_{est} , may be determined by applying the acid balance (eqn. (2)) of the mineral acid. As a rule of thumb, the number N of successive additions S_n and measurements of pH and T to be made before executing the regression analysis to determine S_{tot} , may advantageously be chosen to be about half the selected value for k . Then the risk of obtaining a $S_{n, \text{acc}}$ being larger than S_{tot} becomes very small and at the same time obtain sufficient empirical determinations of the pH and temperature response of the aqueous mineral acid solution to obtain an acceptable representative determination of S_{tot} . There is no need for making each addition, S_n , of fly ash equally large. The method according to the first aspect of the invention will function equally well as long as the accumulated amount does not significantly overshoot the required S_{tot} to reach the selected target pH and as long as each amount S_n being added is measured and accounted for in the accumulated amount $S_{n, \text{acc}}$.

The amount S_n of each successive addition of fly ash may advantageously be made relatively small after having made approximately $\frac{3}{4}$ of the k additions to avoid that the stop criteria; $S_{n, \text{acc}} < S_{\text{tot}}$ does not lead to the final successive addition of fly ash being so large that the $S_{n, \text{acc}}$ significantly overshoots the empirically determined maximum amount S_{tot} . This may in one embodiment be obtained by setting $S_n = \frac{S_{\text{est}}}{ak}$, where a is an integer selected to be from 2 to 10, preferably from 3 to 10, more preferably from 5 to 10, when n becomes equal to or larger than the greatest integer less than or equal to $\frac{3}{4}$ of k , i.e. when $n \geq \lfloor 0.75k \rfloor$. The parenthesis $\lfloor x \rfloor$ as used herein, denotes the floor function which returns the greatest integer less than or equal to a real number x .

The method according to the first aspect may further comprise a refinement of the determination of S_{tot} by performing a new measurement of pH and T and then determining the parameters β_1 , β_2 , γ_1 , and γ_2 and applying these to determine S_{tot} once more after each successive addition larger than N , i.e. which follows after the first determination of S_{tot} in step 5). This may be obtained by adding a step jj) to be executed after step j) and before step jjj) in step 6):

jj) when the added amount S_n is dissolved, measuring and registering the temperature and pH of the aqueous mineral acid solution at this n 'th step as T_n and pH_n , respectively, and determining and registering an accumulated amount of fly ash, $S_{n, \text{acc}}$, being added to the aqueous mineral at this n 'th step, and then repeat steps 4) and 5) to provide a refined determination of S_{tot} .

This refinement provides the advantage that each addition of S_n and following registration of how the pH and T of aqueous mineral acid solution responds to the added fly ash being made until S_{tot} is achieved, are being applied in the regression analysis such that the determination of S_{tot} includes maximum available empirical data over the entire range of accumulated amount of added fly ash. I.e., the estimate of S_{tot} is thus based on real time modelling of all available datasets from the actual process run while ongoing. In this manner, the regulation of the feed of fly ash becomes independent of prior knowledge of the composition of the fly ash being added, optionally also of the concentration of the mineral acid being applied, and adapts automatically to the variations in the fly ash composition by being made according to empirical information on how the temperature and the pH of suspension under formation responds to the addition of the fly ash from beginning to the end of the addition of fly ash. This example embodiment of the regulation is thus well suited to be used when the solid fly ash being applied is e.g. known to have relatively huge variations in the composition such as e.g. MSWI-fly ash. The formation of the suspension may advantageously comprise stirring of the aqueous mineral acid solution during addition of the solid fly ash, by use of an impeller, gas stirring, or in any other known or conceivable manner.

- The aqueous mineral acid solution applied to dissolve dissolvable compounds in the fly ash may advantageously be a strong acid to provide a high capacity for solving the basic compounds in the fly ash. In practice, the mineral acid should have an initial pH of 2 or lower to enable solving sufficient amounts of the (metal containing) basic salts in the fly ash, preferably an initial pH of 1.5 or lower, and most preferably an initial pH of 1.0 or lower. The leaching, i.e. the dissolution rate of fly ash, may be enhanced by adding one or more oxidizing agents to the aqueous mineral acid solution, such as e.g. O_3 , H_2O_2 , or any other known and conceivable oxidizing agent.
- 5 The invention may apply any mineral acid, preferably a mineral acid such as e.g. hydrochloric acid, nitric acid, or other mineral acid. The mineral acid may advantageously be a waste fraction from other industrial processes, such as e.g. hydrochloric acid formed when scrubbing combustion gases from incineration of municipal waste.
- 10 In a second aspect, the invention relates to a dissolution reactor for forming a suspension of partly dissolved fly ash in an aqueous mineral acid solution, comprising:
- 15 a reactor space (1) adapted to receive and contain a mass, L , of the aqueous mineral acid solution at initial concentration, C_0 , and initial pH_0 ,
- 20 a pH-sensor (2) adapted to measure and register the pH in the aqueous mineral acid solution, and optionally a temperature sensor (3) adapted to measure and register the temperature of the aqueous mineral acid solution, and,
- a supply of solid fly ash (4) adapted to feed an amount S_n of solid fly ash into the aqueous mineral acid solution in the dissolution unit to form a suspension at
- 25 a controlled flow rate and to register the accumulated mass, $S_{n, acc}$, of solid fly ash being added,
- a supply (5) of aqueous mineral acid solution adapted to feed the amount L of aqueous mineral acid solution to the reactor space (1), and
- a control unit (6), which is:
- 30 adapted to receive the registered the pH-values from the optional pH-sensor (2), and optionally the registered temperatures from the temperature sensor (3), and
- adapted to regulate the supply of solid fly ash (4) and optionally the supply (5) of aqueous mineral acid solution,
- 35 and which:
- comprises a computer hardware with computer software contains instructions which when executed, utilises the registered pH from the pH sensor (2), and optionally the temperatures from the optional temperature sensor (3), and controls the supply of solid fly

ash (4) to regulate the feed of solid fly ash according to the method of the first aspect of the invention.

The dissolution reactor according to the second aspect of the invention is schematically illustrated in figure 1, and may operate as follows: The reactor space (1) which is adapted to receive and contain a mass L of aqueous mineral acid solution is filled with mineral acid solution from a mineral acid supply (5). The mineral acid supply may advantageously be controlled by the control unit (6), or alternatively be operated manually or by another process control unit. When the reactor space (1) has received the mass L of mineral acid, the control unit (6) engages the supply (4) of solid fly ash to initiate feeding solid fly ash to the aqueous mineral acid solution. The supply (4) measures and registers the mass flow solid fly ash being added to the acid to determine the accumulated mass, S which has been added. This information is passed to the control unit (6). The temperature sensor (3) and optionally the pH-sensor (3) measures and registers the temperature, and optionally the pH of the aqueous mineral acid solution before and at successive intervals of time during the entire feeding of the fly ash, and passes this information to the control unit (6). The control unit (6) applies this information and executes the instructions in its computer software to regulate the feed of solid fly ash to form the suspension. The dissolution reactor according to the second aspect of the invention may advantageously further comprise an impeller (7) for stirring the mineral acid during the formation of the suspension. The operation of the impeller may advantageously be controlled by the control unit (6).

In a third aspect of the invention, the method for extracting one or more metals from solid fly ash, wherein the method comprises:

- 25 selecting a target pH, pH_{targ} , providing an optimized dissolution rate of those salts in the fly ash containing the one or more metals to be extracted and form a suspension according to any of claims 1 to 10,
- passing the formed suspension to a separator separating the suspension into a solid fraction of undissolved fly ash and a first liquid fraction containing dissolved fly ash,
- 30 disposing the solid fraction of undissolved fly ash,
- adding a base to the first liquid fraction to increase the pH to a level where one or more solved metals in the first liquid fraction precipitates as metal hydroxides,
- 35 passing the first liquid fraction with precipitated metal hydroxides to a separator separating the first liquid fraction with precipitated metal hydroxides into a solid fraction of precipitated metal hydroxides, and a second liquid fraction, and
- collecting, and optionally passing the solid fraction of precipitated metal hydroxides to further processing.

The term “selecting a target pH, pH_{targ} , providing an optimized dissolution rate” as used herein, means that the final pH of the suspension to be obtained by the addition of solid fly ash is aimed at providing the maximum dissolution rate of the one or more salts in the solid fly ash containing the one or more metals to be extracted.

- 5 The process steps of separation of the suspension into a liquid fraction and then to separate at least parts of the liquid's content of metal ions by increasing the pH until they precipitate as metal hydroxides followed by separating and collecting the precipitated metal hydroxides, such process steps have been in industrial use and is thus well known to the person skilled in the art.
- 10 The invention according to the third aspect may apply any known or conceivable method for separating solids and liquid of the suspension, such as e.g. filtration. Likewise, the invention may apply any known or conceivable method for separating precipitated solids from a liquid phase. The separation of precipitated metal hydroxides from the liquid residue may be obtained in any known or conceivable
- 15 manner, such as e.g. filtering, centrifuging, settling, gas stirring, etc. The separated and collected precipitated metal hydroxides may be passed to a post-processing and or refining to be converted to commercial commodities. Typical examples of commercial commodities which may be made from the precipitated metal hydroxides include, but are not limited to, elemental metals such as Al, Ca, K, Na,
- 20 Si, Cr, Cu, Pb, Zn, or an oxide, or a hydroxide, or a chloride thereof. Of these, chemical compounds of one or more of the elements Al, Zn, Pb, and Cu are especially preferred commercial commodities of interest for the present invention.

The major part of the heavy metal content of fly ash is solvable in strong acids. The solid fraction separated from the suspension thus consists mainly of remains

25 considered non-hazardous to the environment and may be disposed as ordinary waste in land-fills, or alternatively, it may be applied as raw material in other industrial processes in any known or conceivable manner.

The precipitation of metal hydroxides may advantageously be performed in a precipitation reactor having one or more pH-sensors to enable regulating the precipitation process. The solubility of most metal hydroxides shows a marked

30 dependence on pH and tends toward a minimum with increasing pH [ref 2]. For example, the solubility of hydroxides of non-amphoteric elements such as e.g. Ca and Be have a decreasing solubility with increased pH up to around pH = 8, hydroxides of amphoteric elements such as Al, Cs, Co, Cu, Ni, Pb, Zn, or Sn has a

35 minimum solubility at around pH = 7, and the solubility of oxyanion-forming elements such as e.g. Cr, Mn, Mo, Sb, V, Ag, Se, or W, do not decrease before pH becomes above around pH = 10. This feature enables precipitating tailored compositions of metal hydroxides by adding base to reach one or several predetermined pH-levels. For example, a first metal hydroxide or mixture of metal

hydroxides may be precipitated by adding base to reach a first predetermined pH and then separating and collecting the precipitated hydroxides, and then adding further base to precipitate a second metal hydroxide or mixture of metal hydroxides at a second predetermined pH, etc.

- 5 The invention may advantageously apply a strong base, selected from one or a mixture of the following: NaOH, KOH, Ca(OH)₂, CaO, or Mg(OH)₂, as addition to the first liquid fraction and cause metals to precipitate. The addition of base may advantageously be combined with stirring the liquid residue from the solid-liquid separation by use of an impeller, gas stirring, or in any other known or conceivable
10 manner.

- The second liquid residue usually contains valuable rest-fractions of metals since the solubility of the metal hydroxides are not zero. The method according to the third aspect of the invention may thus further comprise a post-processing of the liquid residue from the separation of the precipitated metal hydroxides, comprising:
15 mixing the second liquid fraction with solid magnetic particles having one or more adsorbents on their surface which selectively adsorbs the metal ions to be extracted,
feeding the mixture of magnetic particles and the second liquid residue to a magnetic separator separating the mixture into a fraction of magnetic particles and a
20 third liquid fraction,
desorbing the adsorbed metal ions from the adsorbents on the magnetic particles and collect the desorbed metal ions as a product fraction,
disposing the third liquid fraction,
and optionally,
25 regenerating the one or more adsorbents on the desorbed magnetic particles.

- The magnetic particles, may advantageously have a d₅₀ particle size in the range of from 10 nm to 10 μm, preferably from 50 nm to 5 μm, and most preferably from 100 nm to 1 μm, and they may either be ferromagnetic, ferrimagnetic, super-paramagnetic or combinations of these properties. As used herein, the term “d₅₀
30 particle size” means the median diameter of the particle size distribution. For example, if D₅₀=10 μm, then 50% of the particles in the sample are larger than 10 μm and 50% are smaller than 10 μm. The particle size distribution may be determined according to standards ASTM E2651-13 or ISO 13317-1-2001.

- The mixing of magnetic particles and liquid may be obtained by stirred tanks, static
35 mixers, tubular reactors or combinations. The magnetic particles may be separated from the liquid by means of appropriate magnetic fields, either set up by permanent magnets or electromagnets in any known or conceivable manner.

The adsorbent on the particle surface depends on which metallic ions that are to be adsorbed. The invention may apply any known or conceivable adsorbent able to

adsorb metallic ions from an aqueous solution. Examples of suited adsorbents include, but is not limited to; polyaspartic acid, polystyrenesulfonate, chitosan, polyacrylamide, or non-coated magnetite or elemental iron particles, and mixtures thereof.

- 5 The particles and adsorbed metal ions are separated by means of one or several known techniques typically including a first step of exposing the magnetic particles with adsorbed metal ions to an environment that facilitates desorption. In principle, this could be an acidic or basic solution, or an organic solvent. In some systems, the contaminants may be exposed to an environment that facilitate reaction with the
10 contaminants, causing the contaminants to release from the adsorbents.

- After desorption, the magnetic particles may advantageously be washed, dried, quality checked and then, if necessary, subject to a surface conditioning process restoring the surface properties of the particles. The surface conditioning may comprise coating new adsorbent etc. Particles that pass the quality control and thus
15 considered to be practically indistinguishable from new and unused particles, may advantageously be transferred to a particle reservoir for later use.

- The third liquid fraction exiting the separator is usually sufficiently stripped of heavy metals to be considered a non-hazardous material, and may be disposed as a non-hazardous waste. However, the third liquid fraction contains often sufficient
20 Cl-ions to be desalinated to form road salt or other products.

- In a fourth aspect of the invention, the dissolution reactor according to the second aspect is implemented in a plant for extracting metals from fly ash, by the dissolution reactor according to the second aspect further comprising:

- 25 a first separator (8) adapted to separate the suspension (9) formed in the dissolution reactor (1) into a solid fraction (10) of undissolved fly ash and a first liquid fraction (11) containing dissolved fly ash,
a precipitation reactor (12) adapted to receive the first liquid fraction (11),
a supply of a base (13) adapted to add base to the first liquid fraction (11) contained in the precipitation reactor (12),
30 a second separator (14) adapted to receive the liquid fraction containing precipitated metal hydroxides (15) and to separate it into a solid fraction (16) of precipitated metal hydroxides and a second liquid fraction (17).

- The plant according to the fourth aspect of the invention is schematically illustrated in figure 2. As seen on the figure, the plant comprises a similar dissolution reactor
35 for producing a suspension (9) as described above for the second aspect of the invention. The formed suspension (9) is passed into a first separator (8) which separates the suspension (9) into a solid fraction (10) consisting of the undissolved remains of the fly ash, and a first liquid fraction (11) consisting of the aqueous mineral acid solution and the dissolved fly ash. The first liquid fraction (11) is

passed on to a precipitation reactor (12) where the pH of the first liquid fraction is increased by adding one or more bases from a base supply (13) to make at least parts of the solved metals to precipitate in the form of solid metal hydroxides. The liquid fraction (15) having precipitated metal hydroxides is then passed on to a
5 second separator (14) which separates the liquid fraction (15) having precipitated metal hydroxides into a solid fraction (16) of precipitated metal hydroxides and a second liquid fraction (17).

As described above for the third aspect of the invention, the separators (8, 14), precipitation reactor (12) and the supply of base (13), are commonly applied process
10 equipment well known to the person skilled in the art. The invention according to the fourth aspect may apply any known or conceivable separator, precipitation reactor.

As described above for the third aspect of the invention, the second liquid fraction (17) exiting the second separator (14) usually contains valuable rest fractions of
15 metal ions. The invention according to the fourth aspect may thus further comprise:

- a supply (19) of magnetic particles having one or more metal ion adsorbing adsorbents on their surface,
- a mixing tank (18) adapted to receive the second fluid fraction (17) and magnetic particles from supply (19) and to mix the liquid and particles,
- 20 a magnetic separator (22) adapted to receive the mixture (21) of the second liquid fraction and magnetic particles, and to separate the mixture (21) into a solid fraction (25) of magnetic particles and a third liquid fraction (24).

The mixing of particles and liquid in mixing tank (18) may advantageously be enhanced by use of an impeller (20).

- 25 The magnetic particles and adsorbents may advantageously be the same as described above for the third aspect of the invention. Likewise, the plant according to the fourth aspect of the invention may further comprise any process equipment for regenerating used magnetic particles (fraction (25)) and pass them into supply (19) for reuse, and/or for extracting the adsorbed metal ions and converting them
30 into commercial commodities as described above.

List of figures

Figure 1 is a schematic drawing of an example embodiment of a dissolution reactor according to the invention.

- 35 Figure 2 is a schematic drawing of an example embodiment of implementation of the dissolution reactor shown in figure 1 into a plant for extracting metal content from fly ash.

Figure 3 is a schematic drawing of the same plant as shown in figure 2, but with mixing reactor and magnetic separator unit for extraction of rest fractions of metal ions.

Verification of the invention

- 5 A series of four laboratory tests were performed to verify the regulation of addition of fly ash according to the first aspect of the invention. Each test applied 3000 ml of concentrated or nearly concentrated hydrochloric acid having an initial pH of from -1 to 0, as the aqueous mineral acid solution and having a temperature around 20 °C (ambient room temperature).
- 10 Then, successive amounts, S_n , of solid MSWI fly ash from the incineration plant for municipal waste at Haraldrud in Oslo, Norway was added to the HCl-acid under stirring. The composition of the fly ash was not determined, i.e. the verification tests were carried out with unknown contents of dissolvable basic salts in the fly ash being added. After each successive addition and mixing of fly ash into the HCl-acid,
- 15 the pH and temperature of the resulting suspension/aqueous mineral acid solution was measured using a Thermo Scientific Eutech pH 450 equipped with both a pH-sensor and a thermometer.

- After five successive additions S_n of fly ash (i.e. $N = 5$) and subsequent measurements of resulting pH and temperature, the empirical data and eqns. (6) and (7)
- 20 were applied to determine the parameters β_1 , β_2 , γ_1 , and γ_2 by a Gauss-Newton algorithm, and the determined parameters β_1 , β_2 , γ_1 , and γ_2 were applied in eqn. (8) to determine S_{tot} . In addition, the determined parameters β_1 and β_2 , were applied in eqn. (6) to determine a target temperature increase, ΔT_{targ} , to be reached when obtaining the selected target pH, i.e. to determine ΔT_{targ} from the relation:

$$25 \quad (10^{-pH_{targ}})^2 = \beta_1 - \beta_2 \Delta T_{targ}$$

Table 1 Results of test 1, selected target pH: 1.3

n	Measured			Modelled					
	$S_{n, acc}$ [g]	pH _n	ΔT_n [°C]	ΔT_{targ} [°C]	S_{tot} [g]	β_1	β_2	γ_1	γ_2
1	61.5	-0.1	1.8						
2	102.5	-0.1	3.2						
3	143.6	0.0	4.6						
4	164.1	0.0	5.4						
5	225.6	0.1	7.9	11.3	301.6	1.946	0.172	82.2	-303.4

6	246.1	0.1	8.6	12.1	322.9	1.891	0.156	84.1	-262.6
7	266.6	0.1	9.3	12.9	345.9	1.840	0.143	86.5	-219.7
8	287.1	0.1	10.0	13.7	370.6	1.788	0.130	88.9	-179.1
9	307.1	0.2	10.8	13.8	375.8	1.785	0.129	90.5	-155.1
10	328.1	0.3	11.6	13.7	376.1	1.792	0.131	91.6	-139.4
11	348.6	0.5	12.1	13.4	375.8	1.812	0.135	94.4	-102.3
12	369.1	0.8	12.6	13.3	377.8	1.827	0.137	97.8	-60.5
13	379.4	1.3	12.2	13.1	382.6	1.845	0.141	103.3	5.9

The determination of S_{tot} and ΔT_{targ} was refined by repeating the determination of the parameters β_1 , β_2 , γ_1 , and γ_2 by the Gauss-Newton algorithm after each additional addition and mixing of S_n until the selected target pH was obtained in the suspension/aqueous mineral acid solution. The results and added amounts, S_n , are summarized in tables 1 to 4 below.

The results given in Table 1 show that the method according to the first aspect of the invention is able to fairly accurate utilize measurements of how the pH and temperature of aqueous mineral acid responds to successive additions of fly ash, and to utilize this empirically determined response to determine a threshold value for how much fly ash to be added in order to obtain a selected target pH (which usually is selected to represent an optimal dissolution degree of one or more metallic salt(s) which is/are to be extracted from the fly ash). The threshold value determined by the method, S_{tot} , is within 1 % from the measured accumulated amount $S_{n, \text{acc}}$ being when the selected target pH, pH_{targ} , is obtained. Thus, even though the pH-measurements are applied as stop criterium in the example embodiment, instead of the accumulated amount $S_{n, \text{acc}}$, the example embodiment show that the method according to the first aspect of the invention provides a method for regulation the feed of fly ash to obtain an optimized dissolution degree of the metal salts to be extracted from the fly ash.

The importance of including the temperature response in the early stage of the fly ash addition, when the pH-measurements are less reliable, is illustrated by the results of the three next example embodiments presented in tables 2, 3 and 4, respectively. In these example embodiments 2, 3 and 4, it was added a relatively large portion of fly ash in the early stage such that the regression analysis was fed with scarce empirical data over the temperature response at the early stage of the addition. As a result, both the modelled temperature ΔT_{targ} and the modelled total amount of fly as, S_{tot} , became slightly overestimated (by 5 to 10 %) as compared to the registered temperature when terminating the addition. This illustrates that to obtain the best fit for the temperature model, datasets should be recorded and

evaluated at relatively equal intervals in S throughout the process. To minimize deviations due to heat loss, it is advantageous to attempt maintaining a relatively constant fly ash feed rate.

- 5 The results from all four tests demonstrates that the measured temperature can be used actively in control algorithms for the leaching of fly ash in acid to support the pH measurements is feasible and preferable.

10 Table 2 Results of test 2, selected target pH: 1.2

n	Measured			Modelled					
	$S_{n, acc}$ [g]	pH _n	ΔT_n [°C]	ΔT_{targ} [°C]	S_{tot} [g]	β_1	β_2	γ_1	γ_2
1	22.8	-0.2	0.5						
2	91.0	-0.1	3.1						
3	136.5	-0.1	4.5						
4	204.8	-0.1	6.8						
5	273.1	0.0	9.3	16.2	458.4	2.355	0.145	95.9	-66.8
6	341.3	0.1	12.2	16.5	445.7	2.344	0.142	91.9	-127.9
7	364.1	0.6	12.6	14.8	415.2	2.426	0.164	94.8	-87.3
8	375.5	0.9	12.7	14.1	405.5	2.466	0.174	97.6	-51.1
9	386.8	1.2	12.6	13.8	404.3	2.490	0.181	101.5	-4.0

Table 3 Results of test 3, selected target pH: 1.2

n	Measured			Modelled					
	$S_{n, acc}$ [g]	pH _n	ΔT_n [°C]	ΔT_{targ} [°C]	S_{tot} [g]	β_1	β_2	γ_1	γ_2
1	21.3	-0.1	0.5						
2	255.9	0.0	8.9						
3	277.3	0.0	9.6						
4	298.6	0.1	10.4						
5	319.9	0.2	11.2	17.8	484.2	1.692	0.095	95.3	-93.6

6	341.2	0.3	11.9	16.4	458.0	1.738	0.106	99.1	-52.8
7	362.6	0.4	12.7	15.6	439.5	1.778	0.114	100.1	-43.1
8	373.2	0.6	13.6	15.2	418.0	1.810	0.119	93.7	-108.1
9	383.9	0.8	13.2	14.8	416.3	1.837	0.124	99.6	-48.7
10	394.6	1.2	13.3	14.5	417.0	1.858	0.128	105.5	8.6

Table 4 Results of test 4, selected target pH; 1.5

n	Measured			Modelled					
	S _{n, acc} [g]	pH _n	ΔT _n [°C]	ΔT _{targ} [°C]	S _{tot} [g]	β ₁	β ₂	γ ₁	γ ₂
1	20.0	0.0	0.8						
2	180.3	0.1	9.6						
3	200.3	0.1	10.7						
4	220.3	0.3	11.9						
5	240.4	0.5	12.9	17.2	311.3	1.119	0.065	99.7	-69.8
6	260.4	1.0	13.6	15.9	302.1	1.156	0.072	109.4	-25.7
7	270.4	1.5	13.8	15.3	298.6	1.178	0.077	115.6	85.8

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