Non-Dispersive Extraction of Ge(IV) from Aqueous Solutions by Cyanex 923: Transport and Modeling Studies

Hossein Kamran Haghighi 1, Mehdi Irannajad 1,*, Maria Teresa Coll 2 and Ana Maria Sastre 3

1 Department of Mining and Metallurgical Engineering, Amirkabir University of Technology, Tehran 1591634311, Iran; h.kamran.h@aut.ac.ir
2 Department of Chemical Engineering, Universitat Politècnica de Catalunya, EPSEVG, Av. Víctor Balaguer s/n, 08800 Vilanova i la Geltrú, Spain; m.teresa.coll@upc.edu
3 Department of Chemical Engineering, Universitat Politècnica de Catalunya, ESTEIB, Av. Diagonal 647, 08028 Barcelona, Spain; ana.maria.sastre@upc.edu

* Correspondence: iranajad@aut.ac.ir; Tel.: +98-21-6454-2937

Received: 27 May 2019; Accepted: 7 June 2019; Published: 11 June 2019

Abstract: Transport process of germanium from an aqueous solution containing oxalic acid and 100 mg/L Ge was studied. Cyanex 923 immobilized in a polytetrafluoroethylene membrane was employed as a carrier in a flat-sheet supported liquid membrane (FSSLM) system. The speciation of the germanium ion in the oxalic acid medium and related diagrams were applied to study the transport of germanium. The effective parameters such as oxalic acid, carrier concentration, and strip reagent composition were evaluated in this study. Based on the experimental data, the oxalic acid and carrier concentrations of 0.075 mol/L and 20% v/v were the condition in which the efficient germanium transport was achieved, respectively. The concentration range of 0.04–0.1 mol/L was selected for sodium hydroxide (NaOH) as a strip reagent providing the best efficiency to transport germanium through the supported liquid membrane (SLM) system. Furthermore, the permeation model was obtained to calculate the mass transfer resistance of the membrane (Δm) and feed (Δf) phases. According to the results, the values of 1 and 1345 s/cm were found for Δm and Δf, respectively.

Keywords: germanium; supported liquid membrane; transport; Cyanex 923; modeling

1. Introduction

Germanium is a strategic metalloid mainly used in a wide range of high-technology devices [1]. Germanium is found in lead-zinc ores [1] and coal fly ashes [2]. From the perspective of hydrometallurgy, the leach solutions obtained from zinc plant residues and coal fly ashes contain germanium along with the other metals. The most important elements that can be found in these solutions are heavy metals such as zinc, nickel, cobalt, cadmium [3,4]. The germanium separation from these solutions is a vital objective in obtaining purified germanium.

Gasification coal fly ashes (GCFAs) containing germanium can be leached with water to dissolve germanium as water-soluble species [5]. Several processes have been developed to recover and separate germanium from impurities such as precipitation [6], flotation [7], ion-exchange [8], distillation [2], adsorption [9], liquid–liquid extraction (LLX) [10], and supported liquid membrane (SLM) [11–13]. Among these techniques, the LLX processes have been extensively applied to separate germanium from aqueous solutions [3,14]. However, some disadvantages such as high loss of extractant, high capital cost, and difficult operation make this method inappropriate for treating low concentrations of ions. The supported liquid membrane techniques have been introduced as alternative methods to
overcome these disadvantages [15]. High selectivity, easy operation, low consumption of carrier, and the one-step process are some advantages of SLMs [16–19].

Liu et al. [1] reported oxalic acid as an efficient reagent to dissolve germanium from zinc residues. Therefore, one of the important leachates containing germanium is oxalate solution. To separate germanium from this type of solution, a few studies have been carried out. Liquid–liquid extraction of germanium from zinc leachates containing oxalic acid solutions has been carried out using tertiary amines [20,21]. Since species of germanium oxalates are in the neutral form (H$_2$Ge(C$_2$O$_4$)$_3$) [14], it was anticipated that a neutral extractant could be used to extract germanium. In this regard, Cyanex 923 (a mixture of four trialkylphosphine oxides, alkyl = normal, C$_6$, C$_8$ [22]) with solvation extraction behavior was selected. Solvent extraction experiments using Cyanex 923 showed good efficiency for the extraction of germanium from oxalate solutions [14]. To overcome the challenges mentioned for an LLX system, an SLM system containing Cyanex 923 was firstly developed to transport germanium from an oxalate solution.

Cyanex 923 has been widely used in SLM systems. The permeation of cadmium (H$_{(n-2)}$(CdCl$_n$)) from wastewater containing chloride anions has been investigated through SLM processes using Cyanex 923 [23]. Chromium(VI) neutral species (H$_2$CrO$_4$) were transported across an FSSLM system using Cyanex 923 from chloride solutions [24]. Zinc(II) species in the form of H$_2$ZnCl$_2$(2n) were permeated by a solid-SLM from chloride medium using Cyanex 923 [25]. In addition, an FSSLM system has been used to transport HFeCl$_4$ species through a polyvinylidene difluoride (PVDF) membrane with the carrier of Cyanex 923 from a solution containing chloride ions [26]. Alguacil et al. [27] investigated the transport of Au(III) in the form HAuCl$_4$ through a PVDF membrane film impregnated in Cyanex 923 from an HCl solution. Results showed good transport efficiency for gold species. Neutral complexes of uranium(VI) (UO$_2$(H$_2$PO$_4$)$_2$) were separated through an SLM system from phosphoric acid solutions using the mobile carriers containing 2-ethyl hexyl phosphoric acid mono-2-ethyl hexyl ester (PC88A) and Cyanex 923 [28].

The presented research describes a flat-sheet supported liquid membrane (FSSLM) process with a Cyanex 923 carrier, in which the facilitated transport of germanium species is carried out from a solution containing oxalic acid. Significant parameters affecting the transport of germanium such as germanium concentration in the feed phase, carrier composition, oxalic acid concentration, membrane type, and NaOH concentration in the strip phase have been investigated in detail. Finally, a mass transfer model was developed to find $\Delta_{aq}$ and $\Delta_{org}$ values which are the resistances corresponding to the species diffusion through the feed–membrane interfacial layer and membrane phase, respectively.

2. Experimental Section

2.1. Materials

Cyanex 923 (93%) consisting of a mixture of four trialkylphosphine oxides (alkyl = normal, C$_6$, C$_8$) was supplied by CYTEC Inc., NJ, USA. Various concentrations of Cyanex 923 (5–30% v/v) as a carrier were prepared by dissolving it in kerosene (Sigma-Aldrich, MO, USA). In all experiments, purified water was supplied through a water purifier (Siemens, Germany). Water solutions used in this study contained 100 mg/L of germanium ion which were prepared by dissolving GeO$_2$ (99.99%) from Sigma-Aldrich. Desired amounts of oxalic acid powder, from Panreac, Barcelona, Spain, were added to the aforementioned solutions to obtain different oxalate concentrations in the range of 0.05–0.2 mol/L. Sodium hydroxide (NaOH) was used to prepare stripping solutions.

Poly tetra fluoro ethylene (PTFE) and polyvinylidene difluoride (PVDF) flat-sheet hydrophobic membranes with the characteristics being porosity of about 85%, 47 mm diameter, pore size of 0.45 μm, and diameter of 47 mm from Millipore, KGaA, Darmstadt, Germany, were used in the present study as a liquid membrane support.
2.2. Membrane Experiments

FSSLM experiments were carried out in a system including two cells attached together with a flanged chamber between the cells in which a membrane filter could be placed. The effective membrane area was calculated to be $11 \text{ cm}^2$. The configuration of this system has been shown as Figure 1. To prepare membranes for experiments, PTFE membranes were immersed in Cyanex 923 solutions for 30 min. The carrier molecules fill membrane pores by capillarity during this time. Afterward, the membrane was taken out from a Cyanex 923 solution and it was washed with water to remove the carrier excess. Finally, the membrane was placed between two cells. The samples, with volumes of 0.5 mL from both cells, were taken to evaluate the concentration of the metal ions during the experiments. The composition of solutions was determined using inductively coupled plasma atomic emission spectroscopy (ICP-AES Agilent, Santa Clara, CA, USA). The reproducibility of experimental values was calculated in the range of ± 5% and illustrated as error bars in the next figures.

![Figure 1](image-url)  
**Figure 1.** Flat-sheet supported liquid membrane (FSSLM) system used in the current study. 1: Motors, 2: agitators, 3: flat-sheet membrane chamber, 4: feed phase, 5: strip phase, 6: Plexiglass body.

2.3. Transport Equations

Transport phenomena in SLM processes principally occur in three steps including the reaction of species with the carrier at the feed–membrane interface, diffusion across the membrane and stripping at the membrane–strip interface. According to the literature, the flux of species ($J$) can be found by disregarding the concentration of germanium in the strip phase with respect to Equation (1) [16]:

$$J = P_f C_f$$

(1)

In this equation, $P_f$ and $C_f$ represent the permeability coefficient at the feed–membrane interface and the ions concentration in the feed phase, respectively. Furthermore, the flux can be written according to Fick’s first law in a differential form as Equation (2):

$$J = -\frac{V}{A} \left( \frac{dC_f}{dt} \right)$$

(2)

where $V$ and $A$ are the volume of the feed phase and the membrane efficient area, respectively. With respect to Equation (1), the integration form of Equation (2) can be written as Equation (3):

$$\ln\left(\frac{C_{f0}}{C_{f,t}}\right) = -\frac{AP_f t}{V}$$

(3)
where \(C_{f,0}\) and \(C_{f,t}\) depict the concentration of ions in the feed phase at the initial time and time of \(t\), respectively. Hence, the permeability coefficient can be evaluated from the slope corresponding to the plot of \(\ln(C_{f,0}/C_{f,t})\) against \(t\). Furthermore, the transport efficiency (\(\%T\)) of germanium is calculated as Equation (4):

\[
\%T = \frac{C_{f,t}}{C_{f,0}} \times 100
\]

3. Solution Chemistry

Germanium dioxide can be dissolved in water. However, this dissolution is slowly carried out as an intermediate germanium tetra-hydroxide (\(\text{Ge(OH)}_4\)). These species can be converted to germanic acid with the solubility product \((K_{sp})\) of \(2.39 \times 10^{-4.5}\) as Equation (5) [29]:

\[
2\text{H}_2\text{O}_{(aq)} + \text{GeO}_2 \rightleftharpoons \text{Ge(OH)}_4_{(aq)} \rightleftharpoons \text{H}_2\text{GeO}_3_{(aq)} + \text{H}_2\text{O}_{(aq)}
\]

Germanium can form various complexes with organic acids such as tartaric acid, citric acid, oxalic acid, etc. Oxalic acid is a possible complexant which forms various complexes with germanium ion. According to the literature and observations, the solubility of germanium increases in the presence of oxalic acid [1]. The formation of germanium anionic and neutral species has been reported. For instance, Pokrovski et al. [30] described that germanium (0.02 mol/L) and oxalic acid (0.1 mol/L) form anionic species of \(\text{Ge(OH)}_2\text{(ox)}_2^{2-}\) (“ox” depicts oxalate) at pHs below 7. Furthermore, Liu et al. [21] introduced \(\text{Ge(ox)}_3^{2-}\) as an anionic complex of germanium and oxalic acid. The concentrations of germanium and oxalic acid in the latter study were reported to be 0.013 and 0.67 mol/L, respectively. However, others [14,29] reported the formation of neutral species of trisoxalato germinates as Equation (6):

\[
3\text{H}_2\text{C}_2\text{O}_4_{(aq)} + \text{GeO}_2_{(s)} \rightleftharpoons \text{H}_2\text{Ge(C}_2\text{O}_4)_3_{(aq)} + 2\text{H}_2\text{O}_{(aq)}
\]

In the current study, germanium neutral species in the form of trisoxalato germinates have been considered as predominant species participated in the reactions. According to the literature review [23,31,32], the probable extraction reaction of germanium using Cyanex 923 can be written as Equation (7) [14]:

\[
2\text{H}^+_{(aq)} + 4\text{L}_{(org)} + \text{C}_2\text{O}_4_{(aq)}^{2-} \rightleftharpoons \text{H}_2\text{Ge(C}_2\text{O}_4)_3\cdot 4\text{L}_{(org)}
\]

4. Results and Discussion

4.1. Determination of Appropriate Oxalic Acid Concentration in the Feed Phase

With respect to the “solution chemistry” section, germanium was transported across FSSLM as a neutral complex (trisoxalato germinate) using Cyanex 923. The transport efficiency depends on the complexion of germanium with oxalate. Therefore, the concentration of oxalic acid is a significant parameter for transport. In this regard, a series of experiments were conducted at the oxalic acid concentration in the range of 0.05–0.2 mol/L with the carrier concentration of 20% v/v. Figure 2a,b illustrates the obtained results. As can be seen in these figures, the transport efficiency and permeation coefficients decrease at lower and higher values of this range. Moreover, at the oxalic concentration of 0.075 mol/L, the maximum permeation coefficient and transport efficiency were found reaching the values of \(9.97 \times 10^{-4}\) cm/s and 88%, respectively. The species of germanium for various oxalic acid concentrations have been illustrated in Figure 3. As can be seen in this figure, at very low oxalic acid concentrations, the dissociation of oxalic acid (\(\text{H}_2\text{ox}\)) is not complete and the germanium tetra-hydroxide (\(\text{Ge(OH)}_4\)) is the predominant compound. However, by increasing the oxalic acid concentration (up to 0.1 mol/L), the concentration of oxalate is enhanced and it promotes an increase in the \(\text{H}_2\text{Ge(C}_2\text{O}_4)_3\) concentration and consequently the germanium transport improves. However, with
more enhancement of oxalic acid concentration, the concentration of germanium oxalate decreases resulting in the decrease in the germanium transport at a higher oxalic acid concentration (>0.1 mol/L).

![Graph showing the effect of oxalic acid concentration on transport and permeability coefficient of germanium on the FSSLM system.](image)

**Figure 2.** The effect of oxalic acid concentration on (a) transport and (b) permeability coefficient of germanium on the FSSLM system ([Cyanex 923] = 20% v/v, temperature of 22 °C, and [NaOH] = 0.1 mol/L).

![Graph showing species of germanium in various oxalic acid concentrations and 100 mg/L Ge.](image)

**Figure 3.** Species of germanium in various oxalic acid concentrations and 100 mg/L Ge (calculated by Medusa software, KTH university, Stockholm, Sweden).
4.2. Evaluation of Appropriate Carrier Concentration

The presence of mobile carriers in a membrane phase is necessary for the transport phenomena; however, an excess amount enhances the viscosity and decreases transport [33]. Therefore, in order to obtain efficient transport, determining an optimum concentration of the carrier is important. In the present study, taking into account the results from solvent extraction experiments, the range of 0–30% v/v for the Cyanex 923 concentration was selected. All FSSLM experiments were done with 0.075 mol/L of oxalic acid, at temperature of 22 °C, and using 0.1 mol/L of NaOH as a receiving solution. According to the result plotted in Figure 4a,b, the complete reaction between Ge(IV) species and carrier molecules does not occur at lower concentrations of the carrier. The germanium transport efficiency rises to about 88% when carrier concentration increases from 10% v/v to 20% v/v. Moreover, with increasing carrier concentration, the curves corresponding to Cyanex 923 of 30 and 25% v/v are shifted down, showing the decrease in germanium transport. In Figure 4b, this behavior can be observed in the reduction of the permeability coefficient. The mentioned reduction may be due to the enhancement of viscosity resulted from the increase in Cyanex 923 concentration and its precipitation in pores of the membrane [34]. According to the aforementioned discussion, the concentration of 20% v/v is selected as the optimum carrier concentration.

![Figure 4. Effect of the carrier concentration on (a) transport and (b) permeability coefficient of germanium on the FSSLM system ([Oxalic acid] = 0.075 mol/L, temperature of 22 °C, and [NaOH] = 0.1 mol/L).](image-url)
4.3. Effect of NaOH Concentration as a Strip Reagent on Germanium Transport

Stripping is an important reaction that takes place in the membrane–strip phase boundary layer on an SLM. For this reaction to occur, the selection of an appropriate striping reagent—a chemical substance such as an acid or a base—is important. To select efficient reagent/reagents for stripping germanium from the loaded organic carrier, a series of liquid–liquid extraction experiments were carried out, and the results have been published elsewhere [14]. Among several reagents such as ammonium chloride (NH$_4$Cl), sodium hydroxide (NaOH), catechol (C$_6$H$_6$O$_2$), citric acid (C$_6$H$_8$O$_7$), ammonia (NH$_3$), sodium sulfate (Na$_2$SO$_4$), and sulfuric acid (H$_2$SO$_4$), NaOH was selected because it is the most efficient reagent for stripping germanium from 20% v/v Cyanex 923. Therefore, FSSLM experiments were carried out to evaluate the effect of the NaOH concentration on the germanium transport using the carrier concentration of 20% v/v Cyanex 923 and an oxalic acid concentration of 0.075 mol/L in the feed solution. The results can be seen in Figure 5a,b.

![Graph showing the effect of NaOH concentration on transport efficiency and permeability coefficient of germanium](image)

**Figure 5.** Effect of NaOH concentration on (a) transport and (b) permeability coefficient of germanium on the FSSLM system ([Oxalic acid] = 0.075 mol/L, temperature of 22 °C, and [Cyanex 923] = 20% v/v).

As can be seen in these figures, the transport efficiency and the permeability coefficient increase with an enhancement of NaOH concentration. About 87% of germanium was transferred to the strip phase at concentrations in the range of 0.04–0.1 mol/L NaOH. The transport efficiency corresponding to the NaOH concentration of 0.02 mol/L is approximately 66%. Furthermore, according to Figure 5b, the permeability coefficient rises to $9.28 \times 10^{-4}$ cm/s at the NaOH concentration of 0.1 mol/L. This is due to the increase in the amount of OH$^-$ anions which enhances the de-complexation rate at the...
interface of the strip side [31]. The probable reaction for stripping species from the loaded carrier can be written as Equation (8) [14]:

$$H_2\text{Ge}(C_2\text{O}_4)_3 \cdot 4L_{\text{org}} + 2OH^- \rightleftharpoons 4L_{\text{org}} + \text{Ge}(C_2\text{O}_4)_3^{2-}_{\text{(aq)}} + 2H_2O_{\text{(org)}}$$  (8)

4.4. Effect of Membrane Type on Germanium Transport

To know the effect of membrane type on the transport of germanium, an experiment was conducted at the condition of 100 mg/L Ge, oxalic acid of 0.075 mol/L, and 0.1 mol/L of NaOH. For this experiment, a polyvinylidene difluoride (PVDF) membrane support was used.

As seen in Figure 6, there is not a significant difference between the transport efficiencies corresponding to the PTFE and PVDF membranes obtained from the experiments carried out under a similar condition. However, the overall transport efficiency of the PTFE membrane is somewhat greater than that of PVDF. The values of the germanium permeability coefficients of the PVDF and PTFE membranes were found to be $3.14 \times 10^{-4}$ and $9.28 \times 10^{-4}$ cm/s, respectively. Therefore, the germanium permeability through the PTFE membrane is approximately three times that through PVDF. Similar results have been reported by Adnan et al. [35]. Various parameters such as higher tortuosity of PVDF membrane result in reducing their permeability [36].

![Figure 6. Transport efficiency of germanium through polytetrafluoroethylene (PTFE) and polyvinylidene difluoride (PVDF) membranes versus time ([Oxalic acid] = 0.075 mol/L, temperature of 22 °C, [Cyanex 923] = 20% v/v, and [NaOH] = 0.1 mol/L).](image)

4.5. Permeability Model

The permeation of germanium ions through the FSSLM (with the PTFE membrane) was modeled to find the mass transfer resistances. To model the mentioned system, it was assumed that species transport through SLM was done with diffusion and the chemical reactions took place instantly [37]. Figure 7 illustrates how germanium oxalate permeates through a PTFE membrane containing Cyanex 923 based on the following steps:

(i) In the feed phase, germanium oxalate and protons diffuse to the interface layer.
(ii) Diffused species and Cyanex 923 molecules react together in the mentioned layer.
(iii) The produced complexes permeate across the membrane toward the membrane–strip interface layer.
NaOH detaches germanium-Cyanex 923 complexes at the membrane–strip phase interface. Therefore, germanium species are stripped from the organic carrier.

The unloaded carrier molecules permeate inversely toward the feed phase.

Figure 7. A schematic transport of germanium ion through FSSLM-Cyanex923.

The extraction reaction of germanium by Cyanex 923 has been described in Equation (7). According to an LLX model developed by Haghighi et al. [38], the extraction equilibrium constant of the aforementioned reaction was calculated to be 2056. The germanium flux can be obtained using Fick’s first diffusion law. Hence, the fluxes at the feed–membrane boundary layer and the membrane phase (\(J_f\) and \(J_m\), respectively) can be presented as Equations (9) and (10):

\[
J_f = \frac{1}{\Delta_f} ([\text{Ge(IV)}]_f - [\text{Ge(IV)}]_{bl,f}) \tag{9}
\]

\[
J_m = \frac{1}{\Delta_m} ([\text{H}_2\text{Ge(ox)}_3\cdot4\text{R}]_{bl,f} - [\text{H}_2\text{Ge(ox)}_3\cdot4\text{R}]_{bl,s}) \tag{10}
\]

where \(\Delta_m\) and \(\Delta_f\) depict the resistances corresponding to the membrane phase and the feed phase boundary layer, respectively. Subscripts \(f\), \(bl\), and \(s\) are feed, boundary layer, and strip, respectively. Moreover, \([\text{Ge(IV)}]_f\), \([\text{H}_2\text{Ge(ox)}_3\cdot4\text{R}]_{bl,f}\), and \([\text{H}_2\text{Ge(ox)}_3\cdot4\text{R}]_{bl,s}\) represent the germanium concentrations in the feed phase, the feed–membrane, and the strip-membrane boundary layers, respectively.

It is noted that since the germanium concentration in the strip-membrane boundary layer is lower than that in the feed–membrane boundary layer, \([\text{H}_2\text{Ge(ox)}_3\cdot4\text{R}]_{bl,s}\) has been neglected. Hence, Equation (10) can be rewritten as Equation (11):

\[
J_m = \frac{1}{\Delta_m} ([\text{H}_2\text{Ge(ox)}_3\cdot4\text{R}]_{bl,f}) \tag{11}
\]

Since the chemical reactions took place instantly, the flux values in the feed–membrane layer and the membrane phase are equal (\(J_f = J_m = J\)). Therefore, overall \(J\) can be found as Equation (12):

\[
J = \frac{K[H^+]^2[R]_{org}^4[\text{Ge(IV)}]_f}{\Delta_m + \Delta_f(K[H^+]_{ay}^2[R]_{org}^4)} \tag{12}
\]
Regarding Equation (1), the permeability coefficient is also written as Equation (13):

$$P = \frac{K[H^+]^2[R]_{\text{org}}^4}{\Delta_m + \Delta_f (K[H^+]^2[L]^4[R]_{\text{org}}^4)}$$  \hspace{1cm} (13)

By arranging Equation (13), Equation (14) allows us to obtain the mass transfer resistances:

$$\frac{1}{P} = \Delta_f + \frac{\Delta_m}{(K[H^+]^2[L]^4[R]_{\text{org}}^4)}$$  \hspace{1cm} (14)

By plotting $\frac{1}{P}$ versus $\Delta_f$, a straight line is obtained.

The intercept and the slope values of this line were used to determine $\Delta_f$ and $\Delta_m$, respectively. This plot is shown in Figure 8. According to the plot, the values of $1$ and $1345 \text{ s/cm}$ were found for $\Delta_m$ and $\Delta_f$, respectively.

**Figure 8.** The plot of $1/P$ vs. $\frac{[H^+]^4}{[K[H^+]^2[L]^4]}$ ([Ge] = 100 mg/L, [Oxalic acid] = 0.075 mol/L, temperature of 22 °C, and [NaOH] = 0.1 mol/L).

5. **Conclusions**

This paper focuses on studying the transport of germanium(IV) through an FSSLM using Cyanex 2932 as a carrier. The main finding of this study is that germanium and oxalate can form neutral species that can be extracted and transported by Cyanex 2932 through an FSSLM system. Results show that the oxalic acid concentration is an essential parameter for the germanium transport because extractable germanium oxalate species can also form in a limited range of oxalate concentration. An optimum concentration of oxalic acid for the complete transport of germanium is in the range of 0.075–0.1 mol/L.

The appropriate Cyanex 2932 concentration in the liquid membrane phase for the complete germanium transport is 20% v/v. Due to the opposite effect of the viscosity at the higher concentrations of Cyanex 2932 on germanium transport, the concentrations over 20% v/v are unsuitable for an effective process. In strip solutions with NaOH concentrations of 0.04–0.1 mol/L, the maximum germanium transport is achieved. Finally, a permeation model was developed to find the mass transfer resistances. As a result, the values of $1$ and $1345 \text{ s/cm}$ were found for $\Delta_m$ and $\Delta_f$, respectively.

**Author Contributions:** H.K.H. made a significant contribution to every stage of this paper, such as the investigation, analysis, methodology and visualization. M.I. contributed to the conceptualization, project administration, validation and supervision of the paper. M.T.C. contributed to the investigation for the paper. A.M.S. contributed to the funding acquisition, investigation and resources provision for the paper. M.I. and H.K.H. finalized the paper by a critical revision.
**Funding:** This research was partially funded by THE SPANISH MINISTRY OF ECONOMY AND COMPETITIVENESS, MINECO, grant number CTM2017-83581-R.

**Acknowledgments:** This research was implemented in the Department of Chemical Engineering, Universitat Politècnica de Catalunya (Barcelona-Tec), Spain. The authors acknowledge Agustin Fortuny for his help. This work has been partially supported by the Spanish Ministry of Economy and Competitiveness, MINECO, through the grant number CTM2017-83581-R.

**Conflicts of Interest:** The authors declare no conflict of interest.

**References**

1. Liu, F.; Liu, Z.; Li, Y.; Wilson, B.P.; Lundström, M. Recovery and separation of gallium(III) and germanium(IV) from zinc refinery residues: Part I: Leaching and iron(III) removal. *Hydrometallurgy* 2017, 169, 564–570. [CrossRef]


5. Torralvo, F.A.; Fernández-Pereira, C. Recovery of germanium from real fly ash leachates by ion-exchange extraction. *Min. Eng.* 2011, 24, 35–41. [CrossRef]


21. Liu, F.; Liu, Z.; Li, Y.; Wilson, B.P.; Liu, Z.; Zeng, L.; Lundström, M. Recovery and separation of gallium(III) and germanium(IV) from zinc refinery residues: Part II: Solvent extraction. *Hydrometallurgy* 2017, 171, 149–156. [CrossRef]


© 2019 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).