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# Preface

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## PREFACE

The 1<sup>st</sup> International Conference on Science and Technology for Sustainable Industry (ICSTSI) is organized as a collaboration between The Institution of Research and Standardization of Industry Banjarbaru (Balai Riset dan Standardisasi Industri Banjarbaru) and The Chemistry Department of Lambung Mangkurat University. The theme of this conference is *"Emerging Science and Technology as a Solution for Global Challenge on Research and Technology Based on Sustainable Resources"*. Sustainability remains a crucial issue regarding how the industrial sectors use technology to explore and utilize resources in the era of industrial revolution 4.0. This revolution has started with the applications of advanced manufacturing and digital information technologies. The world is undergoing a fundamental transformation, which is changing the way we live, work and think. The fourth industrial revolution presents a future filled with exciting opportunities and new challenges and the potential to meet specific requirements during the COVID-19 crisis. Industry 4.0 technologies have the capability of providing better digital solutions for our daily lives during this crisis.

The main goal of this conference is to bring together scientists, academics, and practitioners who work in various fields to exchange their ideas by presenting their research findings. This two-day of conference are certainly the moment of scientific forum that can be used as a means of knowledge exchange to connect participants (in any discipline) with each other, so they can discuss their work, learning from one another, and achieving improvements in results.

In the initial plan, this conference should have been held in Banjarbaru, a city in the province of South Kalimantan, Indonesia on August 6<sup>th</sup>-7<sup>th</sup> 2020. Unfortunately, due to the extraordinary global health issue with the coronavirus disease (Covid-19) that causes the large-scale social restrictions in the framework of the acceleration of Covid-19 handling, ICSTSI was held virtually while maintaining its original schedule. There were about 250 attendees that participated in the conference meeting, and 154 of them were presenting their papers in the plenary sessions. The meeting was done by virtual instead of being postponed because virtual meetings have become the new normal under COVID-19. The virtual meeting was held by Zoom meeting video-conference method. This virtual conference could be used as a tool to speed up existing discussions and timelines among scientists. Each presenter delivered the paper for approximately 7 minutes presentation and followed by 3 minutes discussions and Q&A sessions. The event was appreciated by the participants because it was well-organized and being one of the pioneers in the new normal video-conference trends. The YouTube channel of the opening ceremony ICSTSI 2020 could be found here https://s.id/u1Kmu.

The plenary session in the first day consisted of the presentation by keynote speakers and poster presentations. The parallel session in the second day contained of invited speakers and oral presentations, divided into 6 different topics: material and applied chemistry; wood and non-wood forest products technology; food, cosmetics, and medicine; analysis and method validation industrial, process optimalization technology; biorefinery, bioenergy, and renewable energy and biotechnology; and waste treatment and environmental management. The keynote and invited speakers who had been presented on this conference are listed below.

# Keynote Speakers:

Dr. Ir. Doddy Rahadi, MT (Ministry of Industry, Indonesia) Prof. Dr. Shinso Yokota (Utsunomiya University, Japan) Assoc. Prof. Dr. Azlan Kamari (Universiti Pendidikan Sultan Idris, Malaysia) Prof Dr. Ir. Umar Santoso, M. Sc. (Faculty of Agricultural Technology Gadjah Mada University, Indonesia) Mukhlis Bahrainy (The CEO of Pachira Group) Prof. Dr. Is Fatimah (The Islamic University of Indonesia)

## **Invited Speakers:**

Prof. Rodiansono, Ph.D (Lambung Mangkurat University)
Prof. Gustan Pari (Forest Products Research and Development Center)
Dr. Nazarni Rahmi (The Institution of Research and Standardization of Industry Banjarbaru, Ministry of Industry)
Ir. Lies Indriati (Center for Pulp and Paper, Ministry of Industry)

Dr. Lisman Suryanegara (Center for Research of Biomaterial, Indonesian Institute of Science)

Dr. Aris Mukimin (Center for Industrial Pollution Prevention Technology, Ministry of Industry)

The committee has received 154 papers from both oral and poster presentations, with the participants spread from many countries like Indonesia, Japan and Malaysia. The participants of conference came from various research institutions and universities such as Indonesian Rubber Research Institute, Palembang Institute for Industrial Research and Standardization; Center for Craft and Batik; Bandar Lampung Institute for Industrial Research and Standardization; Sembawa Research Centre; Department of Metallurgical and Materials Engineering; University of Indonesia; Department of Chemical Engineering; Politeknik Negeri Sriwijaya, Indonesia; Department of Agroindustrial Technology, Bogor Agricultural University; Department of Agroindustrial Technology, Lambung Mangkurat University; Department of Chemical Engineering, Engineering Faculty, University of Lambung Mangkurat; Departement of Chemistry, Gadjah Mada University; Wetland-Based Materials Research Center, Lambung Mangkurat University; State Islamic University of Sunan Gunung Djati, Bandung; Center for Ceramic, Bandung; Department of Chemistry, Universitas Palangka Raya; Institution of Research and Standardization of Industry -Padang; Environmental Technic Study Program of Lambung Mangkurat University; Department of Chemistry, Universiti Pendidikan Sultan Idris; Department of Leather Processing Technology, Politeknik ATK Yogyakarta; Faculty of Animal Science, Gadjah Mada University; Banjarbaru Enviroment and Forestry Research Development Institute; Research Division for Natural Product Technology (BPTBA), Indonesian Institute of Sciences (LIPI); Department of Pharmaceutical Chemistry, Gadjah Mada University; Institute of Research and Standardization of Industry, Banjarbaru; Indonesian Swampland Agricultural Research Institute, Banjarbaru; Research Division for Natural Products Technology-Indonesian Institute of Sciences; Faculty of Agriculture, Kyushu University; Faculty of Agriculture, Pattimura University; Accounting, Pamulang University; Center of energy resources technology and chemical industry, BPPT; Chemical Engineering,

Pamulang University. These papers were reviewed by scientific committee and 85 full papers were accepted to be published in IOP Proceeding Conference Series. We would like to thank the participants who have submitted their papers. We are grateful to IOP Publishing as our publication media partner. We also highly appreciate the sponsors who financially supported this event especially to the Industrial Research and Development Council Ministry of Industry, Indonesia.

# LIST OF COMMITTEE

# Steering Committee

- Head of The Institution of Research and Standardization of Industry Banjarbaru
- Head of Research and Development of Industry, Ministry of Industry
- Head of Agro Industry Research Center, Ministry of Industry
- Head of Department of Chemistry, Faculty of Mathematics and Natural Sciences, Lambung Mangkurat University
- Dean of Faculty of Mathematics and Natural Sciences, Lambung Mangkurat University
- Chairman of Indonesia Association of Food Technologist

# **Organizing Committee:**

Person in Charge	:	Ratri Yuli Lestari, S.Hut., M.Env.
Chair	:	Dr. Nazarni Rahmi, S.TP., M.Si
Vice Chair	:	Sunardi, S.Si., M.Sc., Ph.D.
Secretariat	:	Utami Irawati, S.Si., M.E.S., Ph.D; Rais Salim, S.Hut.; Faiza Elisa Hasfianti,S.Hut., Fitri Yuliati, S.Si.
Treasurer	:	Endang Gembirawati, A.Md, Maria Dewi Astuti, S.Si., M.Si.
Event and protocoler	:	Hamlan Ihsan, S.Si; Desi Mustika Amaliyah, S.T., M.M.; M. Listianto Raharjo, S.T.; Drs. Taufiqur Rohman, M.Si.
Plenary session	:	Rodiansono, Ph.D.; Dr. Evy Setiawati, S.Si., M.T.; I Dewa Gede P. P., S.Si.
Equipments and transp	oort	ation: Isa Anshari; Saibatul Hamdi, S.Hut, M.P.; Budi Tri Cahyana, S.T.; Ahmad Budi Junaidi, S.Si., M.Sc.; Dr. Rahmat Yunus, M.Si.
Food and Beverages	:	Rufida, A.Md.; Nadra Khairiah, S.Si, Dahlena Ariyani, S.Si., M.Si.;
		Kholifatu Rosyidah, S.Si., M.Si.
Publication and Docur	ner	ntation: Dwi Harsono, S.Hut.; M. Sobirin; Vemby Danang N.; Dwi Rasy Mujiyanti, S.Si., M.Si.; Noer Komari, S.Si., M.Kes.

Scientific Committee:

# Editor in Chief:

- Prof. Rodiansono, Ph.D. (Chemistry Department, Lambung Mangkurat University)

# **Scientific Committee Members:**

- Prof. Ris. Dr. Muhammad Hanafi (RC Chemistry-Indonesian Institute of Sciences LIPI)
- Prof. Dr. Enos Tangke Arung, S.Hut, M.P. (Faculty of Forestry, Mulawarman University, Samarinda)
- Prof. Dr. Is Fatimah (Department of Chemistry, Islamic University of Indonesia)

- Prof. Iriyanti Fatyasari Nata, S.T., M.T., PhD. (Faculty of Engineering, Lambung Mangkurat University)
- Prof. Rudianto Amirta, S.Hut, M.P. (Faculty of Forestry-Mulawarman University, Samarinda)
- Prof. Dr. Ir. Subyakto, M.Sc. (Center for Research of Biomaterial, Indonesian Institute of Science)
- Prof. Ris. Dr. Ir. Gustan Pari, M.S. (Forest Products Research and Development Center, Ministry of Forestry, Indonesia)
- Dr. Evy Setiawati, S.Si., M.T. (The Institution of Research and Standardization of Industry Banjarbaru)
- Dr. Ragil Widyorini, S.T.M.T. (Faculty of Forestry, Gadjah Mada University)
- Ir. Lies Indriati (Center for Pulp and Paper (BBPK), Ministry of Industry Indonesia)
- Dr. Dede Heri Yuli Yanto (Center for Research of Biomaterial, Indonesian Institute of Sciences-LIPI)
- Sunardi, S.Si., M.Sc., Ph.D. (Chemistry Department, Lambung Mangkurat University)
- Dr. Lisman Suryanegara, S.Si., M.Agr. (Center for Research of Biomaterial, Indonesian Institute of Sciences-LIPI)
- Muthia Elma, ST., M.Sc., PhD. (Chemical Engineering Study Program Engineering Faculty)
- Agung Nugroho, S.TP., M.Sc., Ph.D. (Department of Agro-Industrial Technology, Faculty of Agriculture, Lambung Mangkurat University)
- Dr. Hendra Wijaya, S.Si., M.Si. (Center for Agro-Based Industry-BBIA, Ministry of Industry Indonesia)
- Dr. Nazarni Rahmi, S.TP., M.Si. (The Institution of Research and Standardization of Industry Banjarbaru)
- Dr. Wiwin Tyas Istikowati, S.Hut., M.Sc. (Faculty of Forestry, Lambung Mangkurat University)
- Dr. Ihak Sumardi, S.Hut, M.Si., PhD. (Bandung Institute of Technology)
- Utami Irawati, Ph.D. (Chemistry Department, Lambung Mangkurat University)
- Adi Setyo Purnomo, S.Si., M.Sc., Ph.D. (Department of Chemistry, Faculty of Science and Data Analytics, Institut Teknologi Sepuluh Nopember, Surabaya, Indonesia)
- Dr. Maherawati, S.TP., M.P. (Faculty of Agriculture, Tanjung Pura University, Pontianak, Indonesia)
- Dr. Andriati Ningrum, S.TP, M.Sc. (Faculty of Agriculture Technology, Gadjah Mada University, Yogyakarta, Indonesia)



# CONFERENCE DOCUMENTATION

Figure 1. Conference guests and participants.



Figure 2. Conference opening ceremony.



Figure 3. Conference invited speaker.



Figure 4. Conference committee.



Figure 5. Behind the scenes.

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# Peer review declaration

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# Peer review declaration

All papers published in this volume of IOP Conference Series: Materials Science and Engineering have been peer reviewed through processes administered by the Editors. Reviews were conducted by expert referees to the professional and scientific standards expected of a proceedings journal published by IOP Publishing:

#### Type of peer review: Double-blind •

All papers came through a basic review which includes checking of initial technical criteria (paper scope and similarity checks). The initial technical criteria were checked by the editorial team. Any paper that out of scope was immediately rejected. The accepted papers received were continued to a peer-review process (submission structure, English language usage, and manuscript content) by two professional experts in a related subject area. The identities of both the authors and reviewers were kept hidden. After completing the peer review process, the process was continued by proofreading and layout.

- **Conference submission management system:** • Google forms that connected to email of 1st International Conference on Science and Technology for Sustainable Industry
- Number of submissions received: 154 papers
- Number of submissions sent for review: 120 papers
- Number of submissions accepted: 85 papers
- Acceptance Rate (Number of Submissions Accepted / Number of Submissions Received X 100): 55.19%
- Average number of reviews per paper: two reviews
- Total number of reviewers involved: 23 reviewers
- Any additional info on review process:

The reviewers were asked to answer a number of questions related to the quality of the content (reviewed form). Thus, the review processes were not personal and formalized. The review process for each paper took more than one month. The decision to accept or reject was based on the suggestions of each reviewer. If differences of opinion occurred between two reviewers, the editorial team pointed out one additional reviewer to review. The decision was based on majority votes based on all comments.

Contact person for queries: seminar.brsbb@gmail.com

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Accepted papers received: 04 November 2020 Published online: 31 December 2020

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H Handayani, A Ramadhan, A Cifriadi, N A Kinasih, A F Falaah and D R Maspanger

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# Co (II) desorption from silica gel and mercapto-silica hybrid

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# Co (II) desorption from silica gel and mercapto-silica hybrid

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Abstract. This paper aims to study Co (II) desorption from silica gel (SG) and mercaptosilica hybrids (MSH) by many desorbing agents. SG was prepared from sodium silicate from rice husk ash, whereas MSH was SG modified by organic compounds 3- (trimethoxysilyl) -1propanthiol. The characterization of adsorbents covered by an X-ray diffractometer and an FTIR spectrophotometer. Optimization of contact time for adsorption of Co (II) on HMS and the procedure of desorption agents (H<sub>2</sub>O, HCl, Na<sub>2</sub>EDTA.2H<sub>2</sub>O) at different contact times and concentrations via batch methods. The results showed that the optimum contact time for Co (II) adsorption on MSH was 120 minutes. The most appropriate desorbing agents for SG and MSH were HCl and Na<sub>2</sub>EDTA.2H<sub>2</sub>O. The optimum contact times for Co (II) desorption on SG and MSH were 90 and 120 minutes, respectively. The optimum concentration of HCl for desorbing Co (II) on SG was 0.025 M. While, the optimum concentration of Na<sub>2</sub>EDTA.2H<sub>2</sub>O for Co (II) desorption on MSH was 0.1 M.

## 1. Introduction

Abundant rice husk from the rice mills had been used as a fuel for combustion of red brick, cooking, media hydroponic plants, or burned away. Another application, waste rice husk ash (ASP) has a great potential as the source of silica-based adsorbent because numerous studies showed that the ASP contains sufficiently high points of silica, equivalent to 98% (w/w) [1]. Mujiyanti, et al. [2] reported that the silica content in the rice husk ash from the peat, Banjar Regency, South Kalimantan extended 95.6% (w/w).

Expenditure of the adsorbent adsorbs heavy metals, became a smaller amount of consideration. The adsorbent can be reused through recovery. Natalia [3] has investigated for desorption studies hybrid silica on metal sulfonate with Ca (II) and Cd (II) ions via technique column. The results showed that the result of desorption was Cd (II) ion, 98% with 0.1 M HCl and 100% with 0.05 and 0.1 M Na<sub>2</sub>EDTA.2H<sub>2</sub>O. On the other side, Ca (II) ion desorbed on HCl 0.5 M, amounting to 78.13% and for Na<sub>2</sub>EDTA.2H<sub>2</sub>O, Ca (II) ion desorbed amounting to 88.66%.

Furthermore to adsorbent recovery, desorption can be carried out for the recovery of adsorbed metals. Through desorption studies, we can also find out the adsorption mechanism between the metal and the adsorbent. Mujiyanti (2007) conducted a study of the multi-metal adsorption-desorption of Ag (I), Pb (II), Cr (III), Cu (II), and Ni (II) through a column system in mercapto-silica hybrids (MSH) using three types of eluents, HCl, H<sub>2</sub>O, and Na<sub>2</sub>EDTA.2H<sub>2</sub>O. The desorption with H<sub>2</sub>O % was variety 0.8-31.0% for monometallic solutions and 0.3-24.4% for multi-metal solutions. Meanwhile, the contribution of the adsorption mechanism for the formation of complex bonds with Na<sub>2</sub>EDTA.2H<sub>2</sub>O in

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monometallic solutions alternated from 0.3-29.4% and multi-metal solutions reached from 0.2-14.3%. The desorption process indicated that the adsorption mechanism tends to the formation of hydrogen bonds. This is maintained by the formation of an octahedral aqua complex from the five metal ions in water.

Mujiyanti, et.al [4], stated that Co (II) adsorption on silica gel (SG) optimum at pH 5 and contact time of 75 minutes, while the MSH takes place optimally at pH 3, but the optimum time on MSH could not be determined. The contact time of adsorption was only until 90 minutes in a 50 mL solution of Co (II) 100 mg/L at pH 3 and the amount of adsorbed metal showed a tendency to rise. Therefore, one of the initial steps in this paper was to study the effect of the optimum contact time for Co (II) adsorption by MSH. Furthermore, this paper will observe some aspects of Co (II) desorption from SG and MSH.

# 2. Materials and methods

# 2.1. Materials

The material used was rice husk ash from peat area, South Kalimantan as a source of silica, NaOH (pa Merck), HCl 37% (pa Merck), 3- (trimethoxysilyl) -1-prophantiol (pa Merck), the stock solution of Co (II) 10000 ppm (pa Merck), Na<sub>2</sub>EDTA•2H<sub>2</sub>O (pa Merck).

# 2.2 Determining optimum contact time of Co(II) adsorption

The procedure of determining optimum contact time of Co (II) adsorption on the MSH referred to research that had been conducted by Mujiyanti,et.al [4]. A total of 100 mg of MSH interacted with 50 mL solution of Co (II) 100 mg / L at pH 3. The solution was stirred with a variation of the contact time 30, 60, 90, 120, and 150 minutes, and strained with filter paper. The filtrate was analyzed by Atomic Absorption Spectrophotometry (AAS).

# 2.3 The adsorption of Co (II) on silica gel and mercapto-silica hybrid

The adsorption process referred to research by Mujiyanti,et.al [4] in optimum conditions, pH = 5 and contact time 75 minutes for SG. For MSH accomplished at pH 3 and contact time obtained in determining the optimum contact time adsorption Co (II) at MSH. A total of 1 g of MSH contacted with 50 mL solution of metal ions Co (II) 100 mg/L at the optimum condition. The mixture was filtered then the filtrate was analyzed by AAS. The same treatment was also examined for SG.

# 2.4 The desorption of Co(II) Ion by silica gel and mercapto-silica hybrid

2.4.1 Determining desorption agent of Co(II) desorption. The desorption agent used in this study were H<sub>2</sub>O, HCl 0.1 M and 0.1 M Na<sub>2</sub>EDTA• 2H<sub>2</sub>O. Total of 25 mL desorption agent contained in erlenmeyer and MSH residues that have adsorbed metal ions Co (II), then solution shaken by contact time for 120 minutes. The mixture strained then the filtrate was analyzed by AAS. The same treatment was also examined for SG.

2.4.2 Determining optimum contact time of Co (II) desorption. Desorption was done as in the determination of desorption agent in variation of 30, 60, 90, 120, and 150 minutes.

2.4.3 Determining optimum concentration of desorption agent for Co (II) desorption. Desorption was done as in the determination of desorption agent and contact time, but consuming the suitable agent with a concentration of 0.025; 0.05; 1; 1.5; and 2 M for optimum contact time.

# 3. Results and discussion

# *3.1 Determining optimum contact time of Co(II) adsorption*

Interaction time between adsorbent and the metal ion was one of the factors that play a role in determining the optimum conditions for interaction of metal ions and the equilibrium state of the adsorbent [5]. Figure 1, showed that the number of ions Co (II) adsorbed MSH on contact time 30, 60, 90, 120, and 150 minutes, respectively



Figure 1. The chart of % ion Co(II) adsorbed with the various contact time.

The quantity of Co (II) ions adsorbed was enlarged with an accumulative interval of stirring (30-120 minutes). The extended contact time of MSH and metal ion approved diffusion and annealing process taken more immaculate adsorbate molecules and the amount of Co (II) ions that absorbed increasingly. Adsorption was assessed by the maximum point when the contact time of 120 minutes, in which the percentage of adsorption MSH compared to Co (II) ion was 24.21%. Adsorption equilibrium was characterized by the lack of the concentration of metal ion alterations both on the surface of the adsorbent and metal ion in solution [6], in which the rate of adsorption was equal to the rate of desorption [7]. After passing through 120 minutes of the percentage of Co (II) ions adsorbed tended constantly, even declining. This trend suggested that the active sites on the surface of the adsorbent. This could occur if the interaction between Co (II) metal ions with the active site on MSH was fragile.

### 3.2 The desorption of Co(II) ion by silica gel and mercapto-silica hybrid

3.2.1 Determining desorption agent of Co(II) desorption. Desorption is the opposite of the adsorption process. Desorption is the process of releasing the adsorbate from the adsorbent into a solution [8]. The desorption process can provide information about the adsorption mechanism that plays a role in the adsorption of Co (II) by SG and MSH adsorbents. Co (II) metal ion, which had been adsorbed by the adsorbent, desorbed by the eluents; H<sub>2</sub>O, 0.1 M HCl and 0.1 M Na<sub>2</sub>EDTA• 2H<sub>2</sub>O by batch method. The relationship between percent Co<sup>2+</sup> which wass desorbed with desorption agent types is showed in figure 2.

Distilled water (H<sub>2</sub>O) as a solvent metal ion was attached to the surface of the adsorbent. Desorption result with distilled water did not have a significant effect to release ions Co (II), just 0.76 and 1.47% for SG and MSH. Co(II) metal ion desorbed on surface SG and MSH by HCl solution was successively as much as 103.46 and 54.53%. HCl was a strong acid group that releases H<sup>+</sup> ions to replace metal ions adsorbed by ion exchange process or through hydrogen bonding interactions. Desorption percentage by HCl on SG exceed 100% due to the addition of HCl was expected to dissolve some of the SG, so that Co(II) less reacted with SG. Desorption with Na<sub>2</sub>EDTA•2H<sub>2</sub>O solution indicated that as many as 93.84 and 67.40% of ions Co(II) previously adsorbed by SG and MSH desorbed. The results showed that the adsorption of Co (II) in the adsorbent occurred such as through the formation of a complex between Co(II) with active sites, mercapto (-SH) in the adsorbent.

EDTA is a strong ligand, so when the ability of the ligand adsorbent active group is weaker than the EDTA then the metal ions on the adsorbent will be adsorbed by the formation of a stable complex [3].



Figure 2. The chart of % ion Co(II) desorbed with the various of desorption agent.

From figure 2, showed that the main interaction on SG and MSH was chemisorption. The adsorption mechanism of Co (II) cations on SG adsorbent was the most dominant estimated by ion exchange and /or hydrogen bonding, because the percentage of desorption with HCl equated to other desorbing agents maximum. However, the adsorption mechanism on MSH adsorbent, assumed mostly through the complex formation, an interaction was stronger than ion exchange interactions and the formation of hydrogen bonds.

3.2.2 Determining optimum contact time of Co (II) desorption. Determination of the optimum contact time of desorption aimed to obtain information on how long it takes to achieve optimum Co (II) desorption from the adsorbent. The variation of contact time was from 60 to 180 minutes. This research was using the desorbent agents 25 ml of 0.1 M HCl and 0.1 M Na<sub>2</sub>EDTA•2H<sub>2</sub>O as. The amount of Co (II) depleted for each variation of contact time is shown in figure 3.



Figure 3. The relationship of Co(II) desorpted in various contact time of desorption.

From figure 3 it was shown that the extended contact time of metal ions using 0.1 M HCl and 0.1 M Na<sub>2</sub>EDTA•2H<sub>2</sub>O in SG and MSH, the number of Co (II) ions were separated greatly. However, at a certain moment the addition of time would decrease the percent desorption Co (II) ion which was separated from the adsorbent. Figure 3(A) indicated that the optimum contact time of desorption achieved after SG interacted with HCl for 90 minutes. This was due to the H<sup>+</sup> ion from HCl was strong enough to replace metal ions attached to the adsorbent SG [9]. While the pattern of desorption time

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relationship with the percentage of Co (II) ions was separated on MSH by  $Na_2EDTA \cdot 2H_2O$  showed in figure 3(B). Desorption of MSH continued to rise until it reached a maximum desorption at 120 minutes, which amounted to 98.54%. For the interaction time of 150 and 180 minutes, separate Co (II) ions decreased. According to Khasanah [10] a similar tendency was possible due to a decrease in the surface of the adsorbent that had been saturated by Co-EDTA complex was formed.

3.2.3 Determining the optimum concentration of desorption agent for Co (II) desorption. This research also studied the effect of the concentration of desorption agent compared to Co (II) desorption (variation of the concentration of the agent 0.025 M to 0.2 M). The relationship between the concentration of HCl with Co (II) ions which desorbed on SG can be seen in figure 4(A). by desorption at 97.82 %.



Figure 4. The relationship % ion Co(II) desorpted on various concentration of desorption agent.

According to Natalia [3], one factor was the effectiveness of competition desorption H<sup>+</sup> ions with adsorbed metal, the concentration of the solvent greatly gave affect the effectiveness of desorption. At the appropriate concentration, the ability of H<sup>+</sup> ion to replace the presence of metal ions on the adsorbent being highest. Figure 4(A) showed the increasing of concentration of HCl had lower percentage desorption of Co (II) ions. It was possible that the solvent concentration was too high affecting Co (II) which were separated partially settled down as CoCl<sub>2</sub>. From this research, it was found that the maximum percentage of Co (II) ions was desorbed at a concentration of 0.025 M for HCl as a desorption agent for the Co (II) attached to SG. Figure 4(B) shows that the accumulated concentration. At a concentration of 0.1 M, the optimum amount of Co (II) ions was absorbed at 95.68%. The increase in the desorbed Co (II) ion is proportional to the increase in the concentration of Na2EDTA • 2H2O. But after the concentration had exceeded the optimum, a decrease Co (II) ion was desorbed. This phenomenon was possible due to the partial of deprotonitation Na<sub>2</sub>EDTA•2H<sub>2</sub>O therefore the acid groups ionized carboxyl was less active and caused the formation of the coordination bonding between EDTA and metal ions was not intensive.

## 4. Conclusion

Based on the results of the research, adsorption of Co (II) ion on MSH had reached a maximum point at a contact time of 120 minutes as much as 24.21%. The maximum desorption process of Co (II) ions from SG with HCl and in MSH with Na<sub>2</sub>EDTA•2H<sub>2</sub>O as a desorbing agent. The contact time used in desorption of Co (II) ions on SG by 0.1 M HCl reached a maximum in 90 minutes, as much as 99.27% and MSH by 0.1 M Na<sub>2</sub>EDTA•2H<sub>2</sub>O was 120 minutes, as much as 98.54%. The concentration of 0.025 M HCl as desorbent ion Co (II) in SG which maximally absorbed Co(II) was 97.82% and the concentration of Na<sub>2</sub>EDTA•2H<sub>2</sub>O in the desorption of ion Co(II) from MSH was 95.68%.

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