

University of Belgrade Technical Faculty in Bor



International Mineral Processing & Recycling Conference



Proceedings

Editors: Jovica Sokolović Milan Trumić

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TABLE OF CONTENTS

PLENARY LECTURES	1
L. Guo, Y. Zhao, Q. Ma, G. Tang, C. Jia, C. Li RESEARCH PROGRESS, TRENDS, AND INNOVATIONS OF DEVELOPMENT ON MINING BACKFILL TECHNOLOGY OF UNDERGROUND METALLIFEROUS MINE	3
V.A. Chanturia, V.V. Morozov, G.P. Dvoichenkova, E.L. Chanturia, Yu. A. Podkamenny INNOVATIVE TECHNOLOGY FOR THE RECOVERY OF ABNORMALLY LUMINESCENT DIAMONDS BASED ON THE USE OF LUMINOPHORE-CONTAINING MODIFIERS	23
G. Vujić N. Maoduš, M. Živančev WTE AS INTEGRATED PART OF CIRCULAR ECONOMY	32
J.C. Gabriel, H. Bo, N. Charpentier, S. Chevrier, Y. Deng, F.Olivier, D. Xia CRITICAL METALS RECOVERY FROM E-WASTE: FROM MICROFLUIDICS HYDROMETALLURGY TO ECONOMICALLY VIABLE PROCESSES	39
SESSION LECTURES	41
F. Nakhaei, I. Jovanović 3D IMAGING AND APPLICATIONS IN MINERAL PROCESSING	43
D. Singh, S. Basu, B. Mishra. R. Bhima Rao NOVEL APPROACHES TO RECOVER TOTAL HEAVY MINERALS FROM DIFFERENT GRADE BEACH SAND DEPOSITS USING GRAVITY CONCENTRATORS	54
M. Trumić, K. Balanović ROLE OF PARTICLE SHAPE IN THE FLOATABILITY OF TONER PARTICLE	64
I. Smičiklas, M. Egerić, M. Jović COPPER SORPTION CAPACITY OF THE SOIL TREATED WITH UNCONVENTIONAL ALKALIZING AGENTS	73
V. Conić, I. Jovanović COPPER ORE BIOLEACHING FROM ECOLOGICAL POINT OF VIEW	79
S. Cvetković, M. Popović, J. Perendija LIFE CYCLE ASSESSMENT AND USE OF NATURAL RESOURCES	89
WORKSHOP PAPERS	95
P. M. Angelopoulos, G. Anastassakis, N. Kountouris, N. Koukoulis, M. Taxiarchou COMBINED USE OF ORGANOSOLV LIGNIN AND XANTHATES ON SPHALERITE FLOTATION FROM MIXED SULPHIDES	97
P. M. Angelopoulos, N. Kountouris, G. Anastassakis, M. Taxiarchou PARTIAL REPLACEMENT OF XANTHATE BY ORGANOSOLV LIGNIN ON PYRITE/ARSENOPYRITE FLOTATION	103
K. Hrůzová, July Ann Bazar, Leonidas Matsakas, Anders Sand, Ulrika Rova, Paul Christakopoulos ORGANOSOLV LIGNIN PARTICLES: A NOVEL GREEN REAGENT THAT INCREASES THE FLOTATION EFFICIENCY OF SULFIDE ORES	109
A. Peppas, D. Skenderas, P.M. Angelopoulos, C. Politi ENVIRONMENTAL BENEFITS OF LIGNIN BASED ECOFRIENDLY SURFACTANTS FOR FLOTATION PROCESSES TOWARDS CURRENT PRACTICES	115

A. Peppas, K. Hurzova, D. Skenderas, C. Politi, L. Matsakas, P.M. Angelopoulos EVALUATION OF BATTERY MINERALS FLOTATION PROCESS ECO FRIENDLINESS UTILISING BIODEGRADABLE LIGNIN REAGENTS	121
A. Peppas, C. Politi, D. Skenderas, P.M. Angelopoulos ENVIRONMENTAL ASSESSMENT OF RARE EARTHS RECOVERY METHOD FROM BAUXITE RESIDUES	126
PAPERS	133
A. Jankovic, M. Sederkennya MODIFIED BOND AND RITTINGER ENERGY-SIZE RELATIONSHIPS FOR LABORATORY FINE GRINDING	135
V. Nikolić, M. Trumić, D. Tanikić OPTIMIZATION OF MICRONIZING ZEOLITE GRINDING USING ARTIFICIAL NEURAL NETWORKS	143
E. Petrakis, K. Komnitsas THE EFFECT OF MICROWAVE RADIATION ON DRY GRINDING KINETICS OF BAUXITE ORE	150
M.H. Tyeb, S. Mishra, A.K. Majumder LSTM AND CNN COMBINATION BASED MODELLING APPROACH FOR PARTITION CURVE PREDICTION IN HYDROCYCLONES	157
I. Jovanović, M.Ž. Trumić, J. Sokolović, M.S. Trumić, J. Nešković DETERMINATION OF LIMITING SETTLING VELOCITY IN THE SLURY PIPELINE FROM GRINDING PLANT, USING DIFFERENT APPROACHES – A CASE STUDY	163
N. Omarova, R. Sherembayeva, A.Amirkhan, Zh. Ibraybekov, A. Nesipbay FLOTATION OF POLYMETALLIC LEAD-ZINC ORES OF THE BAKALSKOYE DEPOSIT	168
V.A. Chanturiya, I.Zh. Bunin, M.V. Ryazantseva THE APPLICATION OF THE DIELECTRIC BARRIER DISCHARGE (DBD) FOR THE IMPROVEMENT OF THE SEPARATION OF PYRITE AND ARSENOPYRITE	174
V. Ignatkina, A. Kayumov, N. Yergesheva, P. Chernova BASIC SELECTIVE REAGENT REGIMES FOR COMPLEX SULFIDE ORE FLOTATION	179
S. Chaudhuri, S. Maity, S.C. Maji, D. Roy, U.S. Chattopadhyay STUDIES ON THE FLOATABILITY CHARACTERISTICS OF LOW VOLATILE COKING COAL FINES USING X-RAY DIFFRACTION (XRD) ANALYSIS AS A DIAGNOSTIC TOOL	186
V.I. Ryaboi, V.P. Kretov, E.D. Schepeta, I.V. Ryaboi, S.E. Levkovets APPLICATION OF COLLECTOR BTF-15221 IN FLOTATION OF COPPER- AND GOLD - CONTAINING ORES	193
I. Dervišević, A. Dervišević, M. Tomović, J. Galjak COMPARATIVE ANALYSIS OF REAGENTS FOR GOLD EXTRACTION FROM FLOTATION TAILS	202
E.M.S. Silva, A.C. Silva, J.M.B.S. Cabral, P.S. Oliveira, A.F. Nascimento, A.P. Vieira Filho, S.A. Santos TESTS WITH DIFFERENT FLOCCULANTS FOR CHROMIUM ORE TAILINGS	208
C. Ouyang, B. Lv, K. Jia, Y. Yang STUDY ON THE APPLICATION OF HIGH-EFFICIENCY AND ENVIRONMENT-FRIENDLY COPPER COLLECTOR TO ASSOCIATED COPPER IN AN IRON ORE	214
S. Sredić, Lj.Tankosić KINETIC STUDIES OF THE ADSORPTION POLYACRILAMIDE-BASED FLOCCULANTS ON NATURAL GOETHITE, QUARTZ AND CLAY MINERALS	221

G. D. Bogdanović, D. Marilović, B. Nikolić, S. J. Petrović COLUMN LEACHING OF LOW-GRADE COPPER SULFIDE ORE WITH SULFURIC ACID	230
K. Gáborová, M. Achimovičová, M. Hegedüs, O. Šestinová AN INFLUENCE OF MECHANICAL ACTIVATION ON THE COPPER LEACHING KINETICS OF BERZELIANITE	236
D. Medić, I. Đorđević, M. Nujkić, A. Papludis, V. Nedelkovski, S. Alagić, S. Milić USE OF COPPER POWDER AS A REDUCING AGENT IN THE LEACHING PROCESS OF LiCoO ₂	242
J. Dimitrijević, S. Jevtić, A. Marinković, M. Simić, M. Koprivica, J. Petrović REMOVAL OF HEAVY METAL IONS FROM MULTIMETALLIC SOLUTION BY MODIFIED OAT STRAW	248
M.R. Rath, A.S. Patra, S. Kiran Kumar, M. Mukherjee, A. Chatterjee, A. Ranjan, A.K. Bhatnagar, A.K. Mukherjee A PROCESS TO DECREASE THE CLAY COATING OF IRON ORE LUMPS & FINES BY THE APPLICATION OF DISPERSANTS	254
H. Kurama, S. Kurama SURFACTANTS AND THEIR FUNCTIONS ON NANO-POWDER SYNTHESIS	262
A. Goryachev, D. Makarov METHODS FOR PROCESSING NATURAL AND ANTHROPOGENIC COPPER- NICKEL RAW MATERIALS IN THE ARCTIC	275
Y. Yuankun, D. Mirović DAM BREACH ANALYSIS USING HEC-RAS: A CASE STUDY OF COPPER AND GOLD "ČUKARU PEKI" MINE DAMS	283
A. Milovanović Brkić, Y. Yuankun, N. Buđelan MANAGEMENT OF FLOTATION TAILINGS AS MINING WASTE ON THE COPPER AND GOLD MINE "CUKARU PEKI"	289
N. Pavlovic, F. Palkovits, A. Hall GEO-STABLE DISPOSAL OF COAL COMBUSTION BYPRODUCTS	297
N. Pavlovic, F. Palkovits, A. Hall TAIL WAGGING THE DOG-WHY INTEGRATED SOLUTIONS ARE BETTER-TAILINGS AND BACKFILL DISPOSAL	303
V. Alivojvodic, N. Petrovnijevic POSITION OF COPPER WITHIN URBAN MINING - RECOVERING POTENTIAL FROM MINE TAILINGS	309
V.Tsitsishvili, N.Dolaberidze, N.Mirdzveli, M.Nijaradze, Z.Amiridze, B.Khutsishvili BACTERIOSTATIC ACTIVITY OF GEORGIAN HEULANDITE ENRICHED WITH BIOLOGICALLY ACTIVE METALS	315
V.Tsitsishvili, M.Panayotova, N.Dolaberidze, N.Mirdzveli, M.Nijaradze, Z.Amiridze, B.Khutsishvili, N.Jakipbekova, S.Sakibayeva THERMAL STABILITY OF NATURAL HEULANDITE-CHABAZITE MIXTURES	321
V.Tsitsishvili, M.Panayotova, N.Dolaberidze, N.Mirdzveli, M.Nijaradze, Z.Amiridze, B.Khutsishvili, N.Klarjeishvili, N.Jakipbekova COMPOSITION OF GEORGIAN AND KAZAKHSTANI NATURAL HEULANDITES	327
S. Matijašević, S. Grujić , V. Topalović, J. Stojanović, J. Nikolić , V. Savić, S. Zildžović NANOCRYSTALLIZATION OF POTASSIUM NIOBIUM GERMANATE GLASSES	333

A.C. Silva, E.M.S. Silva, P.S. Oliveira, A.F. Nascimento, A.P. Vieira Filho, D.B. Carvalho Neto ESTIMATING THE ACCURACY, PRECISION, AND RECALL OF THE HAND-SORTING OF	338
A BRAZILIAN CHROMIUM ORE	
V.V. Morozov, Y.P. Morozov, G. Zorigt, D. Lodoy, E. Jargalsaikhan, I.V. Pestriak SCANNING FLATBED OPTICAL ORE QUALITY ANALYZER	344
B. B. Tchouffa, N. J. Ndemou, M. G. Frida Ntsama CHARACTERIZATION, ENRICHMENT TEST AND VALORIZATION OF IRON ORE FROM NABEBA (NORTH – CONGO)	350
K. Jia, S. Đorđević, C. Ouyang, B. Lv LABORATORY BENEFICIATION TECHNOLOGY AND DEVELOPMENT RESEARCH ON TITANIUM MAGNETITE ORE	355
D. S. Radulović, V. Jovanović, B. Ivošević, D. Todorović, S. Milićević, M. Marković INVESTIGATION OF THE POSSIBILITY OF VALORIZATION OF TWO BORATE SAMPLES FROM THE DEPOSIT "POBRĐE" – BALJEVAC	361
S. Hredzák, M. Matik, O. Šestinová, A. Zubrik, D. Kupka, S. Dolinská, I. Znamenáčková, M. Sisol, M. Marcin, L. Pašek STUDY OF ORE SAMPLES FROM THE ZLATÉ HORY DEPOSIT (HRUBÝ JESENÍK Mts., SILESIA, CZECH REPUBLIC)	367
J. Sokolović, I. Ilić, D. Krstić COMPARISON OF THE RESULTS OF SEPARATION OF DIFFERENT COALS IN THE ANTHRACITE MINE "VRSKA CUKA"	373
B.R. Reddy, K. Abhishek, J.M. Korath, M.R Rath A COMPUTATIONAL TOOL FOR PREDICTION OF JIG CONCENTRATOR OPERATING PARAMETER TO GET IMPROVED YIELD OF CONCENTRATE	379
I. Jovanović, V. Conić, D. Milanović, F. Nakhaei, S. Krstić RELATIVE PREDICTION ERROR OF FLOTATION INDICES BY ANFIS MODELS	387
Z. Štirbanović, R. Stanojlović, J. Sokolović, D. Stanujkić, N. Ćirić, I. Miljanović, G. Popović APPLICATION OF VIKOR METHOD FOR SELECTION OF COLLECTOR IN PORPHYRY COPPER ORE FLOTATION	391
S. Milutinović, Lj. Obradović, S. Petrović S. Magdalinović, I. Svrkota RANKING OF FLOTATION TAILINGS POND IN EASTERN SERBIA USING THE AHP METHOD	398
I. Jovanović, V. Conić, J. Sokolović, D. Kržanović, D. Radulović SIMPLE FUZZY MODELS FOR PREDICTION OF FLOTATION INDICES	404
S. Mishra, M.H. Tyeb, A.K. Majumder DEVELOPMENT OF A VIBRATION SENSOR-BASED ONLINE MONITORING SYSTEM FOR DETECTING ROPING IN HYDROCYCLONES	410
B. Farkaš, A. Hrastov, E. Orbanić THE IMPROVEMENT OF MINERAL PROCESSING – CASE STUDY	416
T. Mohit, P. Patel, P. Kaushal, J. Sahoo, V. Arumuru, B. Deo, M. Jain, R. Manchanda IMPROVED ON-LINE FAILURE PREDICTION METHOD OF COAL INJECTION SYSTEM USED IN A SPONGE IRON ROTARY KILN	423
M. Mikić, R. Rajković, S. Trujić, D. Kržanović, M. Jovanović IMPACT ON THE ENVIRONMENT AND OF THE OPEN MINE AND LANDFILLS IN SOUTH MINING DISTRICT – MAJDANPEK	429

M. Jovanović, D. Kržanović, R. Rajković, M. Mikić, M. Maksimović APPLICATION OF GEOGRIDS IN RECULTIVATION MEASURES AGAINST LAND DEGRADATION	435
V. Gardić, R. Marković, Z. Stevanović, A. Isvoran, T. Marković APPLICATION OF SUSTAINABLE CYCLING MANAGEMENT SYSTEM IN PHYTOREMEDIATION TECHNOLOGY OF CONTAMINATED SOILS	441
D. Đurđević-Milošević, A. Petrović, J. Elez, G. Gagula, V. Kalaba SUSTAINABLE APPROACH TO THE LACTIC ACID PRODUCTION AND ANTIBACTERIAL USE	445
B. Cekova, M. Matlievska, M. M. Puncheva, V. Velkoski, B. Kuzmanovski DIGITALIZATION OF WASTE, WAYS FOR MORE EFFICIENT WASTE MANAGEMENT	451
A. Vasileiadou, S. Zoras, A. Dimoudi INVESTIGATION OF SLAGGING CHARACTERISTICS OF INDUSTRIAL SOLID WASTES	458
A. Vasileiadou, S. Zoras, A. Dimoudi MODELLING OF COX AND NOX EMISSIONS FROM INDUSTRIAL SOLID WASTES COMBUSTION USING ANSYS CHEMKIN PRO	464
Z. Bayer Ozturk, S. Kurama, A. Eser THE USAGE AND EFFECT OF BASALT CUTTING WASTE (BCW) IN CERAMIC GLAZE COMPOSITIONS CONTAINING OPAQUE AND MATT FRIT	470
D. Dinić, S. Stupar, N. Jovanović, M. Tanić, S. Jevtić SYNTHESIS AND CHARACTERIZATION OF POROUS CERAMICS BASED ON COPPER SLAG	480
M. Šišić, Dž. Dautbegović, M. Duraković ANALYSIS OF THE CHARACTERISTICS OF SLAG FROM METALLURGICAL PLANTS IN ZENICA DISPOSED OF INDUSTRIAL WASTE LANDFILL "RACA"	486
Dz. Datubegovic, M. Hasanbasic, M. Sisic, V. Birdahic ANALYSIS OF THE IMPACT OF THE INTRODUCTION OF LARGER CONTAINERS INTO THE WASTE COLLECTION SYSTEM IN THE CITY OF ZENICA	492
N. Bušatlić, I. Bušatlić, A. Halilović, N. Merdić, L. Kovač ENVIRONMENTALLY ACCEPTABLE CEMENTS WITH THE ADDITION OF GRANULATED BLAST FURNACE SLAG	498
A. Stojićević, M. Antić, M. Purić VEGETABLE INDUSTRY BY-PRODUCTS AS RAW MATERIALS IN FUNCTIONAL FOOD PRODUCTION	507
A. Petrović, R. Marković, D. Božić CARBON NANOTUBES AS POTENTIAL MATERIAL FOR WASTEWATER TREATMENT - A REVIEW	514
M. Marić, A. Ivković, B. Ivković, A. Janošević Ležaić, S. Uskoković-Marković, J. Savić, M. Milojević-Rakić, D. Bajuk-Bogdanović REMOVAL OF METHYLENE BLUE FROM AQUEOUS SOLUTIONS USING AN IRON- RICH SOIL	519
R. Marković, V. Gardić, R. Kovačević, Zoran Stevanović, A. Isvoran, V. Marjanović, A. Petrović BOR DISCRICT RIVERS WATERCOURSES CONTAMINATION BY Cu AND NI IONS	524
P. Kekarjawlekar, N. Kamal, K. Maniyar, B. Deo, P. Nanda, P. Malakar, R. Manchanda DEVELOPING SAFE OPERATING PRACTICES (SOP) FOR POSTCOMBUSTION CHAMBER IN A SPONGE IRON PLANT	530

D. Milošević, M. Radosavljević, S. Polavder, Ž. Praštalo ARRANGEMENT OF FIELDS DEVASTATED BY CONSTRUCTION OF MAIN GAS PIPELINE	536
D. Đurđević-Milošević, A. Petrović, J. Elez, V. Kalaba, G. Gagula ENVIRONMENTAL PROTECTION THROUGH THE RATIONAL USE OF SODIUM HYPOCHLORITE AS A FUNGICIDE	542
G. Kyparissis, A. Goulkoudis, G. Papadimas, E. Tasiopoulos, A. Vasileiadou CASE STUDY OF ENERGY SAVING IN A PUBLIC SCHOOL THROUGH THE INSTALLATION OF A PHOTOVOLTAIC SYSTEM ON THE ROOF	548
D. Topalović, J. Marković, M. Jović, S. Dragović, I. Smičiklas THE ARSENIC SORPTION CAPACITY OF DIFFERENT SERBIAN SOILS	554
F. Popescu, M. Zot, E.A. Laza USING SHERPA TOOL FOR ASSESSMENT OF EUROPEAN WATERBORNE TRANSPORT SECTOR IMPACT ON AIR QUALITY	560
A. Stojić, D. Tanikić, E. Požega THE IMPACT OF EXPLOITATION OF PRIMARY AND ALTERNATIVE ENERGY SOURCES ON THE ENVIRONMENT	566
A. Radojević, S. Šerbula, T. Kalinović, J. Milosavljević, J. Kalinović MOBILE PHONES – A VALUABLE COMPONENT OF E-WASTE STREAM	572
K. Janković, M. Stojanović, D. Bojović, A. Terzić, S. Stanković APPLICATION OF COAL COMBUSTION BYPRODUCTS IN SELF-COMPACTING CONCRETE: INFLUENCE ON FLOWABILITY	579
D. Radosavljević, A. Jelić, M. Stamenović IMPACT OF STUDENT MIGRATIONS ON SUSTAINABLE AND TECHNOLOGICAL DEVELOPMENTS OF THE REPUBLIC OF SERBIA	585
D. Radosavljević, A. Jelić, M. Stamenović DEVELOPMENT OF EDUCATION FOR SUSTAINABLE DEVELOPMENT AND MANAGEMENT OF RECYCLABLE WASTE IN THE REPUBLIC OF SERBIA	592
Deependra Singh SUSTAINABLE RECOVERY OF INDIAN PLACER MINERALS-THEIR DISTRIBUTION AND MINERAL ASSEMBLAGES	598
ABSTRACTS	607
M. Tasić, I. Stojković, V. Pavićević, V. Veljković SIMULATION OF HYDRODYNAMIC CAVITATION-ASSISTED BIODIESEL PRODUCTION FROM WASTE COOKING OIL USING ASPEN PLUS	609
A. Jocić, S. Marić, A. Dimitrijević RECOVERY OF METALS FROM INDUSTRIAL EFFLUENTS USING AN IONIC LIQUID- BASED STRATEGY	610
S. Marić, A. Jocić, A. Dimitrijević IONIC LIQUID-BASED TECHNOLOGY FOR METAL RECOVERY FROM ELECTRONIC WASTE	611
J. Vučićević, S. Čupić, M. Jauković, V. Đurđević, M. Stamenović, A. Božić, A. Janićijević CURRENT STATE OF THE QUALITY OF THE LUG RIVER IN THE MUNICIPALITY OF MLADENOVAC	612

XV International Mineral Processing and Recycling Conference, 17-19 May 2023, Belgrade, Serbia	XV International Mineral Proce	essing and Recycling Conference,	, 17-19 May 2023, Belgrade, Serbia
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D. Žnidarič THE ENERGY CRISIS AND THE EXPLOITATION OF MINERAL RESOURCES IN THE	613
LIGHT OF INCREASING LOADS IN SPACE	010
S. Zeković	
A NEW GLOBAL CHALLENGES AND REGULATION FOR SUSTAINABLE SPATIAL	614
DEVELOPMENT OF MINING	
P.M. Angelopoulos, P. Oustadakis, G. Anastassakis, M. Georgiou, N. Kountouris HYDROTHERMAL TREATMENT OF BAUXITE RESIDUE FOR IRON RECOVERY ENHANCEMENT BY MAGNETIC SEPARATION	615
O. Ayoglu, M. Sinche-Gonzalez, M. Moilanen	
TEXTURAL MINERALOGICAL UNDERSTANDING OF MAGNETITE LIBERATION	616
CONTAINING COPPER INCLUSIONS	
M. Sinche-Gonzalez	
MASTER IN MINERAL PROCESING (EMJM-PROMISE) IN THE CONTEXT OF DEMAND	617
OF CRITICAL MATERIALS AND ENERGY TRANSITION	
ADVERTISING MATERIALS	619
Department for Mineral and Recycling Technologies	621
Serbia Zijin Mining	624
Serbia Zijin Copper	627
Analysis d.o.o.	629
tozero	631
Monicom	632
EMJM-PROMISE	633



INVESTIGATION OF THE POSSIBILITY OF VALORIZATION OF TWO BORATE SAMPLES FROM THE DEPOSIT "POBRĐE" – BALJEVAC

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ABSTRACT – Boron is a rare element from the group of non-metals and it occurs in nature in over 250 boron-bearing minerals-borates. Turkey has the largest borate reserves and is their largest producer. Serbia has significant borate resources, for now without exploitation and processing. This paper presents the results of investigating two borate samples from the deposit "Pobrđe. The following properties were determined on borate samples: free and hygroscopic moisture, granulometric composition, chemical composition, mineralogical analysis by size classes, and XRD analysis.

Keywords: "Pobrđe" Deposit, Boron Minerals, Properties.

INTRODUCTION

Boron is a rare element from the group of non-metals (concentration of 10 ppm in the upper continental crust) [1,2]. Boron occurs in nature in over 250 boron-bearing minerals-borates, which are defined as any compound containing boric oxide (B_2O_3). The most important boron minerals in commercial terms are: borax(tincal), kernite, ulexite, colemanite [3]. Approximately 75% of the world's boron reserves are located in Turkey, which is also the largest producer of borates (Table 1) [2].

Countries	Total Reserve (Thousand ton B ₂ O ₃)	Distribution (%)	Productions of borate concentrates, t (2019.)			
Turkey	948,712	73.4	4,000,000			
Russia	100,000	80,000				
USA	80,000	1,300,000				
Peru	22,000	111,108				
Argentina	9,000	0.7	80,000			
China	36,000 2.8		250,000			
Bolivia	19,000	1.5	214,500			
Chile	41,000 3.2		352,225			
Kazakhstan	15,000	1.2	500,000			
Serbia	21,000	1.6	/			
TOTAL	1,291,712	100				

Table 1 World reserves and production of borates [4,5]

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Starting in 2014, Boron was marked as "critical" in the European Commission's assessment study for the E.U. [6,7]. Four main sectors of boron mineral application represent 86% of its world annual consumption: 1. Glass sector -1.17 Mt B₂O₃; 2. Ceramics sector -280,000 B₂O₃ 3. Agriculture sector 240,000 B₂O₃ 4. Detergents sector -45,000 B₂O₃ [8,9]. There are several borate deposits in Serbia: Piskanja - 11.8 Mt [10], Pobrđe - about 150,000 t [11] and it is counted on the possible production of boron from the jadarite mineral [9].

Boron ore from the "Pobrđe" deposit was exploited by "Ibar Coal Mines". After mining, the borates ore was crushed and then sieved on a screen with an opening of 30 mm. Commercial borate concentrate was obtained from the size class +30 mm by hand picking. The size class -30+0.00 mm was discarded to the landfill as tailings. "Elixir group", the largest producer of mineral fertilizers in the Balkans, bought 7-8,000 tons of this raw material and transferred it to the "Elixir Zorka" factory in Šabac. Here, after grinding, this raw material is used as an additive for mineral fertilizer. The "Elixir group" wanted to examine the possibility of obtaining boron mineral concentrates both from run-of-mine ore and from the -30+0.00 mm class. The goal of these tests would be to obtain K/B₂O₃ with a B₂O₃ content of over 35% and a middling product that would have a B₂O₃ content of 7-10% for the fertilizer industry. In order to determine the possibility of conducting magnetic separation tests, detailed physical-chemical and mineralogical characterization was performed.

EXPERIMENTAL (MATERIALS AND METHODS)

Two representative borate samples were taken for these investigations. The run-ofmine borate ore sample had an upper size of limit -280+0.00 mm, mass m=130 kg, and the -30+0.00 mm class sample had a mass of m= 82 kg. On both samples, free moisture was determined. After that, both samples were crushed to a size of -10+0.00 mm, and physico-chemical and mineralogical properties of these samples were determined. The following were determined on the samples: hygroscopic moisture, granulometric composition, chemical composition, mineralogical analysis by size classes, and XRD analysis. Mineralogical analysis by size classes was performed on a polarizing microscope "JENAPOL-U", Carl Zeiss-Jena, with a measuring device. X-ray diffraction (XRD) analysis was performed on a "PHILIPS" X-ray diffractometer, model PW-1710.

Determination of moisture content and granulometric composition of borate ore samples

The content of free moisture in the run-of-mine sample was 2.39% and hygroscopic 0.21%; in the sample of the size class -30+0.00 mm the content of free moisture was 2.51% and the hygroscopic 0.22%.

The granulometric composition for both samples was determined by sieving on a Tyler series of sieves, with the last sieve having an aperture of 0.1 mm. All the weights of the sieves oversize together with the sieve undersize of the last sieve were measured. Granulometric analysis showed that: - for the run-of-mine sample of the borate ore, after three-stage comminution, the d_{95} is 6.895 mm, while the d_{50} is 2.323 mm; - for a borate

sample of the size class -30+0.00 mm, after comminution, d_{95} is 6.773 mm, while d_{50} is 2.193 mm.

Chemical analysis of the borate samples

The chemical composition of both borate samples was determined by chemical analysis. In both samples, the content of the B_2O_3 , and secondary components was determined, before all Fe_2O_3 . Loss of ignition at 800 °C is shown. The chemical compositions of borate samples are shown in Tables 3 and 4.

Table 2 Chemical composition of the run-of-mine borate ore										
Comp.	B ₂ O ₃	SiO ₂	AI_2O_3	Fe ₂ O ₃	CaO	MgO	Na ₂ O	K ₂ O	TiO ₂	LOI at 800°C
Cont.,%	32.21	15.20	4.05	1.61	20.30	3.75	1.06	0.57	0.168	21.03

Table 3 Chemical composition of the borate, size class -30.0+0.00 mm	
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Comp.	B ₂ O ₃	SiO ₂	AI_2O_3	Fe ₂ O ₃	CaO	MgO	Na ₂ O	K ₂ O	TiO ₂	LOI at 800°C
Cont.,%	21.05	21.60	6.09	2.38	20.13	4.83	1.16	0.83	0.33	23.26

Chemical analysis showed that these two borate samples are significantly different from each other. Namely, the starting sample of the borate ore has a much higher boron content (B_2O_3 content 32.21%), compared to the -30+0.00 mm size class sample (B_2O_3 content 21.05%). The starting sample also has a lower content of Fe₂O₃ -1.6% and sample of the size class -30+0.00 mm, has content of Fe₂O₃ -2.38%.

Qualitative-quantitative mineralogical investigation of borate samples

Quantitative-qualitative mineralogical analysis was performed using an optical polarization microscope and X-ray diffraction analysis on several different samples of borate ore. Mineralogical analysis by size class was carried out on crushed borate samples with a size of -10 mm, in order to determine intergrowth and liberations of minerals in each class of the borate samples. These data made it possible to define the size of the class in which boron minerals can be separated from tailings minerals. X-ray diffraction (XRD) analysis was performed on both borate samples after micronization, on the basis of which the mineral composition of the samples was determined.

Mineralogical analysis by size classes of run-of-mine borate sample and borate sample size class -30+0.00 mm

Both borate samples were sieved into the following narrow size classes: -10.00+9.52 mm; -9.52 +7.93 mm; -7.93 + 6.35 mm; -6.35 + 5.00 mm; -5.00 + 4.00 mm; -4.00 + 3.36 mm; -3.36 + 2.83 mm; -2.83 + 2.38 mm; -2.38 +1.6 mm; -1.6 + 1.19 mm; -1.19+0.00 mm. Mineralogical analysis by size classes for both borate samples was performed by examination with an optical polarization microscope. This analysis, for both borate samples, determined that the liberations of boron minerals of over 80% are achieved in the size class -2.83+0.00 mm. Mineralogical analysis by size class of the borate sample of

class -30+0.00 mm showed that this sample has a very dirty surface. The dust stuck to its mineral surfaces, despite crushing and dry sieving.

Mineralogical analysis by X-ray diffraction method

Mineralogical, X-ray diffraction analysis was used to determine and monitor the phase composition of both borate samples. The samples were analyzed on a "PHILIPS" X-ray diffractometer, model PW-1710, with a curved graphite monochromator and a scintillation counter. The intensities of diffracted CuK α X-rays (λ =1.54178Å) were measured at room temperature in intervals of 0.02° 2 θ and time of 1 s in the range from 4 to 65 °2 θ . The X-ray tube was loaded with a voltage of 40 kV and a current of 30 mA, while the slits for directing the primary and diffracted beams were 1° and 0.1 mm. The following text shows the obtained results.

The run-of-mine borate sample - X-ray diffraction method

X-ray analysis determined the following mineral composition of the examined sample: colemanite, howlite, quartz, carbonate minerals (calcite, dolomite), clay minerals (smectite)/chlorites, analcime, jarandolite, and weddellite. The absolutely dominant mineral in the analyzed sample is colemanite. After colemanite, quartz is more significantly present, as well as carbonate minerals (dolomite is more dominant than calcite), clay minerals (smectite), analcime, howlite, and jarandolite are less abundant. The semiquantitative share of crystalline phases (minerals) is as follows: colemanite 55-57%, carbonates 18-20% (dolomite 15-17%, calcite \leq 2-3%), smectites/chlorites ~ 10%, analcime 8-10%, quartz 1-2%, other boron minerals (howlite and jarandolite in total ~ 5%). Weddelite is present in the trace. The diffractogram of the tested sample is presented in Figure 1.

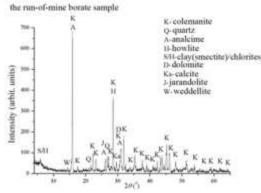


Figure 1 Powder diffractogram of the run-of-mine borate sample

Borate sample of size class -30+0.00 mm - X-ray diffraction method

X-ray analysis determined the following mineral composition of the examined sample: colemanite, howlite, quartz, carbonate minerals (calcite, dolomite), clay minerals (smectite)/chlorites, analcime, and jarandolite.

The absolutely dominant mineral in the analyzed sample is colemanite. After colemanite, quartz is more significantly present, as well as carbonate minerals (dolomite is more dominant than calcite), clay minerals (smectite), analcime, howlite, and jarandolite are less abundant. The semiquantitative share of crystalline phases (minerals) is as follows: colemanite 35-37%, carbonates 30-32 (dolomite 20-22, calcite \leq 10%), smectites/chlorites 10-12%, analcime 8-10%, quartz \leq 5%., other boron minerals (howlite and jarandolite in total ~ 5%). The diffractogram of the tested sample is presented in Figure 2.

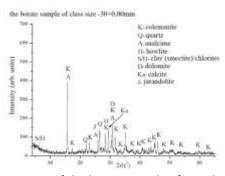


Figure 2 Diffractogram of the borate sample of size class -30+0.0 mm

RESULTS AND DISCUSSION

Physico-chemical and mineralogical investigations represent the basis for further technological tests that should define the processing procedure of both types of borates. These investigations showed that borates are very complex in terms of their mineral composition. Namely, in addition to boron minerals, they also consist of alumino-silicate tailings, oxide carbonate minerals, zeolite (of the analcime type), and organic matter. Based on the mineralogical analysis, the same minerals are present in both borate samples, Figure 1, Figure 2. According to the mineralogical analysis, the content of boron minerals is higher in the run-of-mine borate sample (B_2O_3 -55-57%, Figure 1), than in the borate sample of the size class -30+0.00 mm (B₂O₃- 35-37%, Figure 2).In order to examine and determine the possibility of the separation of minerals one from another, it is necessary that the minerals have some different properties by which they can be separated from each other. Regarding their physical-mechanical properties, hardness by Mohs scale, specific mass, etc. the minerals present in the sample have similar properties. According to the specific mass (density), all these minerals have a specific mass of about 2.5 g/cm³, except montmorillonite, whose specific mass ranges from 2-2.7 g/cm³. Chamosite has a higher specific mass than the mentioned minerals with a specific mass of 3.2 g/cm³. This fact tells us that it would be difficult to separate these minerals by gravity concentration because they are close in density (specific mass).

CONCLUSION

According to the chemical compositions of the minerals present in the borate samples, it can be stated that some of these minerals contain iron, especially smectites,

and chlorites. These minerals can be separated from boron minerals using high-intensity magnetic separation. In addition, montmorillonite itself does not contain iron, but in nature very often binds it to itself, either between lamellae, or by isomorphic replacement, or sometimes both ways. For this reason, montmorillonite very often belongs to the minerals that can be separated by magnetic separation from boron minerals. In any case, in accordance with the mineralogical tests by size classes, the mineral raw material for this purpose should be comminute to a size of -2.83 mm, when the boron mineral is free from accompanying minerals. And after this phase, it is necessary to sieve the raw material prepared in this way into the appropriate size classes that can be subjected to magnetic separation. Through tests, it is necessary to determine the most optimal induction of the magnetic field, during which the magnetic separation of boron and accompanying minerals by size classes that can be subjected to magnetic separation. With these tests, it is necessary to determine the optimal size range of classes that can be subjected to magnetic separation, as well as the last class that cannot be subjected to dry magnetic separation due to its fineness.

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