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ORIGIN AND EVOLUTION OF THE OPHIOLITIC COMPLEX OF ZDRALJICA (CENTRAL SERBIA)

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ABSTRACT

The ophiolitic complex of Ždraljica (Central Serbia) belongs to the Eastern Branch of the Vardar suture zone. It was emplaced during the Upper Jurassic. The complex consists predominately of a MORB/VAB-like tholeiitic suite, represented mostly by gabbros and diabases. Small occurrences of cummulitic peridotites, basalts and plagiogranites also appear. The tholeiitic suite is intruded by calc-alkaline intermediate and acid magmas. Geochemical data suggest that the ŽOC tholeiitic rocks originated by partial melting of a spinel-lherzolite source. Non-modal batch melting modeling indicates 10 to 15 % of partial melting of such a source. The magmas were later modified by fractional crystallization. One-step major element modeling requires 40% (F=0.60) of fractional crystallization of a mineral assemblage: Pl_{52.9}Cpx_{12.5}Ol_{26.1}Ttn_{2.9}Ap_{4.4}Mgt_{1.0}. The model is supported by the variation patterns of most trace elements.

1 INTRODUCTION

The Ždraljica Ophiolitic Complex (ŽOC) is situated near Kragujevac (Central Serbia), about 150 km south of Belgrade. Geotectonically, it belongs to the Eastern Branch of the Vardar Zone Composite Terrane - VZCT (Karamata et al. 1994, Resimić-Šarić et al. 2000), marking, with other similar occurrences, the suture zone of the Vardar ocean and boundary between the VZCT and the Serbo-Macedonian Massive (Fig. 1). The continuation of ophiolitic occurrences can be followed further to the south in Southern Serbia, F.Y.R. of Macedonia and Greece (e.g. Bébien 1977). Northwards, this zone continues as the Mures ophiolites in the Apuseni Mountains in Romania (Zacher & Lupu 1999 and references therein).

The ŽOC covers about 30 km², representing middle and upper parts of the oceanic crust. It mostly consists of gabbros and diabases. The complex is associated with the Upper Jurassic diabase-chert formation (Marković et al. 1968, Dolić et al. 1981) and it is covered by Tithonian limestones and Lower Cretaceous paraflysch sediments as an over-step sequence (Karamata et al. 1999).

The ŽOC started to be investigated in detail few years ago and the first results and interpretation about the evolution of tholeiitic series rocks will be given in this paper. Special attention will be given to processes in the source, magma modification processes and geotectonic setting of the complex. It is argued that the tholeiitic series originated by a moderate degree of partial melting of a lherzolitic upper mantle source. Crystal fractionation was recognized as a main process of magma modification and present major and trace element modeling. Available data suggest that the ŽOC formed in a MORB-like setting, although VA setting cannot be entirely excluded.

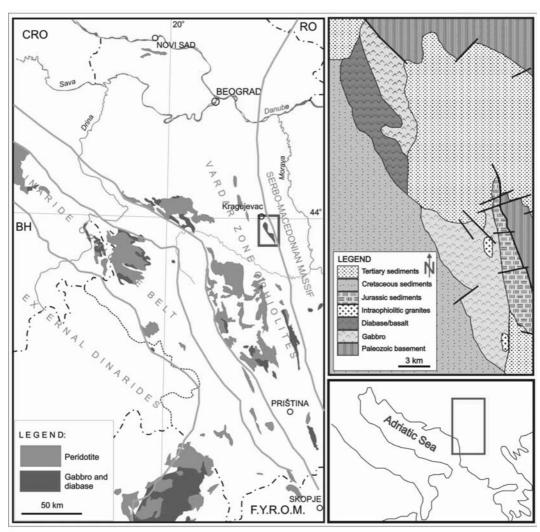


Figure 1. Geotectonic position of the Ždraljica Ophiolitic Complex (ŽOC). The terrane boundaries and distribution of Mesozoic ophiolites are from Karamata et al. (1994) and Dimitrijević (1992). The simplified sketch (upper right) shows geology of the ŽOC after Marković et al. (1968) and Dolić et al. (1981).

2 ANALYTICAL PROCEDURE

Analyses of major and selected trace elements of whole rock samples were done by XRF method at the Department of Earth Sciences of the University of Perugia (Italy). The analyses have been performed using an X-ray tube completed with a Rn and W anode under conditions within a range of 40-45 mV I=30-35 mA. A LiF-200 crystal analizator and scincilating detector were used. Precision is better than 10% for V, Cr, Ni, Y, Zr and Ba, and better than 5% for all the other elements. The accuracy has been tested on international standards, and is better than 10%.

The mineral phases of the rocks were analyzed by Energy Dispersive Spectrometry (EDS) at the Aristotle University of Thessaloniki, Greece. Mineral phases were imaged with backscattered electrons (BSE) and quantitatively analyzed using a LINK AN 1000 EDS microanalyser attached to a JEOL JSM-840 Scanning Electron Microscope. Operating conditions were 15kV accelerating potential, beam current 3nA, surface electron beam $1\mu m^2$ and counting time 80 seconds. The ZAF-4/FLS software provided by LINK was used for corrections. Minerals (albite, orthoclase, diopside, wollastonite, olivine, periclase) and pure metals were used as standards.

The samples were analyzed for K/Ar in the Institute for Nuclear Research ATOMKI of Hungarian Academy of Sciences in Debrecen (Hungary). The ages are reported in Resimić-Šarić et al. (2002) while the analytical procedure is given in Balogh (1985).

3 ROCK CLASSIFICATION AND PETROGRAPHY

Igneous rocks occurring in the ŽOC are generally divided into two genetically different suites: (1) suite of a tholeitic and (2) suite of a calc-alkaline affinity (Resimić 2000). The first suite consists of serpentinised peridotites, cummulitic and massive gabbros, diabases and plagiogranites. The second group comprises intermediate and acid rocks (quartzdiorites, quartzmonzodiorites, granodiorites and leucocratic granites), and they cut the rocks of the tholeitic suite. This paper is focused on the tholeitic rocks of the ŽOC. A detailed study of the calc-alkaline magmatism intruding ophiolites is in progress and these data will be published in a subsequent paper.

Generally, the rocks of the tholeiitic suite show effects of extensive alteration and low- to medium-T metamorphic processes. Therefore, they have sometimes been classified as spilites and metabasalts, or even chlorite-epidote-albite schists and serpentinites. In the following text we will present petrography and mineral chemistry of most fresh igneous rocks of the ŽOC.

Wherlites are developed in the lower part of the Ždraljica ophiolitic complex. They occur as very rare and small cummulitic pods and layers within gabbros. They display allotriomorphic and orthocummulitic texture. The main mineral phases are olivine (Fo_{85.73}) and diopside (Mg# 89.8) and spinels as accessories.

Gabbros occur as irregular blocks or sheets of massive and cummulitic facies. They show coarse-grained, rarely fine-grained and poikilitic, coarse-grained ophitic and orthocummulitic texture. The main constituents are plagioclase (63.8-97.6 %An), sometimes albitized (3.7-19.6 %An), clinopyroxenes (diopside – Mg# 73.0-93.3 and augite – Mg# 62.4-86.7) and, only rarely, totally serpentinized olivine. Accessories are magnetite, sphene and apatite. Secondary minerals are epidote, uralite, chlorite, tremolite, calcite, serpentine, pyrite and rarely magnesiohornblende.

Diabases occur as single dykes or dyke-swarms intruding gabbros or as irregular massive bodies. They show ophitic, intergranular or intersertal texture, sometimes with glassy chilled margins. Plagioclase (70.2-85.7 %An) and augite (Mg# 61.3-80.2) are the main mineral phases, while accessories are magnetite, sphene and glass. Metamorphosed diabases are composed mostly of albite (2.3-6.9 %An), epidote/zoisite, uralite, tremolite, magnesiohornblende, calcite and pyrite.

Basalts appear as pillow lavas, primary and redeposited hyaloclastites and rarely as relicts of coherent lava flows. They show glomeroporphyritic, ophitic, aphyric and intersertal texture. Phenocrysts are ubiquitously albitized plagioclase (7.7-11.7 %An), relicts of augite (Mg# 60.2 and 77.7) and severely serpentinized olivine, while the groundmass is composed of chloritised glass, plagioclase microlites and opaque minerals.

Plagiogranites occur as small dyke- or sill-like intrusions and irregular bodies within diabases. They display hypidiomorphic and fine-grained granular texture. Albite and quartz are the main minerals, accessories are magnesiohornblende, magnetite and apatite, while epidote/zoisite and chlorite are secondary products.

4 AGE OF THE ŽOC

Radiometric age determinations of ŽOC rocks were published in Resimić-Šarić et al. (2002) and here we shall present a brief summary. Previously, the ŽOC was mapped as a part of the diabase-chert formation, the age of which has been variously determined, from Archean to Tertiary (Ćirić 1955, Gočanin 1938, Terzić 1962, Marković et al. 1968).

The presence of an Upper Jurassic over-step sequence proves that the emplacement of the ŽOC occurred before the Upper Jurassic. It is in agreement with the presumed closure of the Eastern Branch of the Vardar Zone (Karamata et al. 1999) and the emplacement of Mures ophiolites (e.g. Bortolotti et al. 2002).

K/Ar analyses on various mineral and whole rock samples of both ŽOC tholeiitic and calcalkaline suites range from 168.4 Ma to 77.3 Ma (Fig. 2). The oldest radiometric age (168.4±6.7 Ma – magnesiohornblende from calc-alkaline quartzdiorite) may correspond to the crystallization age or

a minimum age of the ŽOC ophiolite emplacement. The latter is in agreement with the age of the overstep sequence. To better constrain the significance of these radiometric ages we have to know more about the origin and evolution of the calc-alkaline granitoid rocks. On the other hand, all younger ages (152±21 – 77.3±3.6 Ma) may comprise processes of ocean floor metamorphism or post-emplacement events.

Similar K/Ar ages have been reported on the ophiolitic rocks of the Bükk Mts., N Hungary; 166 ± 8 Ma has been measured on hornblende from gabbro at Szarvaskő and muscovite from the contact of the gabbro resulted 165 ± 5 Ma (Árva-Sós et al. 1987). On the same muscovite an Ar/Ar plateau age of 162.9 ± 0.9 Ma has been published by Balogh & Pécskay (2001). In the SE foreland of the Bükk Mts. borehole Hejőszalonta-1 reached spilitized basalt at 1650.3 m, its age was determined with K/Ar isochron method as 160.6 ± 11.6 Ma and 159.4 ± 3.2 Ma (Balogh & Pécskay 2001).

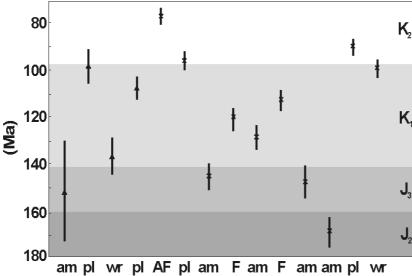


Figure 2. K/Ar data for analysed rocks of the Ždraljica ophiolitic complex (Resimić-Šarić et al. 2002). Explanation: \blacktriangle - gabbros and diabases, \varkappa - calc-alkaline suite; Abbreviations: am – amphibole, pl – plagioclase, AF – alkali feldspar, F – feldspars, wr – whole rock, J_2 – Middle Jurassic, J_3 – Upper Jurassic, K_1 – Lower Cretaceous, K_2 – Upper Cretaceous.

5 GEOCHEMISTRY

Analyses of major element oxides and selected trace elements of 14 samples of basic and one sample of acid tholeiitic rocks are given in table 1.

Their SiO₂ contents range from 30.82 wt.% to 51.31 wt.% while the silica content of plagiogranite is 71.57%. Chemical behavior of major and trace elements is described using MgO as differentiation index. The contents of TiO₂, CaO, FeO_t and P₂O₅ decrease with decreasing of MgO concentrations. SiO₂ and Al₂O₃ contents increase from the most primitive sample to the most evolved one, while contents of Na₂O are rather constant. K₂O contents are scattered. Concentrations of V, Co, Cu, Zn and Ga decrease with decreasing MgO contents while Y and Nb concentrations slightly decrease. Pb and Th contents increase with some scattering, Sr, Ba, Cr and Ni contents are highly scattered while Zr and Ce concentracions do not show changes in their contents during the evolution.

6 DISCUSSION

Analyses of nine selected samples of basic rocks will be included in the discussion. The samples were screened by taking into consideration effects of alteration and metamorphism.

The tholeiitic suite rocks of the ŽOC were plotted on a MORB-normalized (Pearce 1983) spider diagram (Fig. 3). Generally, three groups of rocks can be distinguished: (1) cummulitic gabbros, (2) primitive and evolved gabbros and diabases and (3) a plagiogranite sample.

All the basic rocks show subparallel trace element pattern with cummulitic samples having lower concetrations of most trace elements. The least evolved gabbros and diabases have similar concentrations of most trace elements to MORB. N-MORB affinity of the ŽOC mafic rocks is also inffered from HFSE ratios, e.g. Zr/Y(around 2.5) and Ti/Y (around 240) (Saunders & Turney 1984). In addition, according to (Y+Nb) vs. Rb and Y vs. Nb (Pearce et al. 1984) discrimination diagrams, the ŽOC plagiogranite shows an ORG affinity.

Table 1. Major (wt%) and trace (ppm) element data of rocks of the ŽOC.

Rock-		(NE/8) and trace (ppm) dismont data of reside of the 200.													
type	р	g										d	b	pl-g	
Sam-	1074	1/045		11100		1/0.071			4007	4.000				F504	
ple	V371a	V315	4	M102	8B	V367b	2A	3B	A607	A602	8A	2F	E536d	E521a	E509
SiO ₂	30.82	47.54	50.75	49.03	49.85	46.14	39.45	41.67	49.64	49.64	51.31	39.88	44.07	38.65	71.57
TiO ₂	0.11	1.03	0.53	0.86	0.76	1.05	1.10	0.92	0.28	0.62	1.09	2.21	1.25	2.23	0.42
Al ₂ O ₃	3.13	15.37	15.05	15.76	15.93	22.26	8.52	11.44	15.57	15.75	14.34	8.16	14.61	9.41	12.48
FeO _t	16.39	9.63	11.87	8.76	9.31	5.04	17.18	16.13	7.66	9,85	10.05	18.74	12.98	19.16	5.12
MnO	0.39	0.2	0.14	0.18	0.18	0.11	0.39	0.32	0.15	0.18	0.18	0.44	0.25	0.33	0.11
MgO	34.38	9.71	8.76	8.90	8.71	8.98	8.40	8.06	11.4	8.21	8.46	10.63	9.54	6.92	0.86
CaO	2.56	11.76	13.61	11.57	10.21	11.34	18.05	16.52	11.02	10.19	9.87	11.48	13.88	14.4	2.00
Na₂O	0.07	2.15	1.14	1.95	1.75	1.31	2.28	1.93	1.4	1.86	2.08	4.33	1.35	2.97	5.61
K ₂ O	0.03	0.12	0.09	0.11	0.16	0.08	1.31	0.84	0.08	0.12	0.09	0.72	0.03	0.15	0.03
P ₂ O ₅	0.04	0.27	0.15	0.23	0.17	0.19	0.18	0.16	0.17	0.21	0.19	0.21	0.22	0.28	0.08
LOI	10.37	1.69	2.08	2.16	2.75	4.01	1.90	0.86	2.23	2.73	2.03	1.73		3.84	1.19
V	46	273	247	263	215	50	224	212	155	208	282	279	334	262	43
Cr	2032	451	443	381	358	112	673	603	564	351	244	157	188	115	13
Со	159	45	39	47	38	35	61	68	42	41	42	49	51	68	10
Ni	1813	125	156	112	83	171	228	200	153	100	88	70	62	78	30
Cu	29	73	37	61	74	68	128	123	61	19	15	144	72	81	52
Zn	107	60	39	47	50	27	87	85	36	35	51	129	76	136	45
Ga	4	18	10	14	12	10	24	17	11	13	16	29	16	25	14
Rb	1	1	1	1	1	1	24	19	3	1	1	6	1	1	2
Sr	21	159	265	197	129	156	482	336	142	150	119	207	79	147	77
Υ	6	22	12	25	18	4	25	18	10	17	24	37	25	55	133
Zr	13	73	26	67	48	16	47	35	15	39	71	117	29	130	624
Nb	3	5	0,1	6	3	0,1	3	4	2	3	3	8	4	9	17
Ва	2	8	4	8	4	2	60	29	2	2	10	66	2	21	7
La	0	4	6	4	6	3	13	16	5	7	0	6	13	20	10
Pb	23	7	14	5	9	9	27	20	5	9	8	26	7	22	12
Ce	3	3	2	3	2	3	2	2	3	3	2	2	3	3	89
Th	6	0	1	1	3	3	11	1	2	0	3	5	1	7	4

 $FeO_{t}: total \ iron \ as \ ferrous, \ Abbreviations: p-peridotite, \ g-gabbro, \ d-diabase, \ b-basalt, \ pl-g-plagiogranite$

6.1 Modeling of partial melting and fractionation processes

In order to further elaborate the source and low-pressure modification processes, major and trace element modeling was performed.

Firstly, the Cr vs. Y log-log diagram was used to distinguish rocks, which may represent primary mantle-derived melts from those representing magmas that underwent processes of fractionation (Fig. 4). The non-modal equilibrium (batch) melting equation was used in order to calculate the Cr and Y concentrations for range of 0-40% melting (F=0.0-0.4) of a spinel lherzolite with modal mineralogy $Ol_{57}Opx_{25.5}Cpx_{15}Sp_{2.5}$. Computer iterations were carried out for every 1% melting-step, by recalculation of the mineralogical composition of the source, according to the method proposed by Kostopoulos & James (1992).

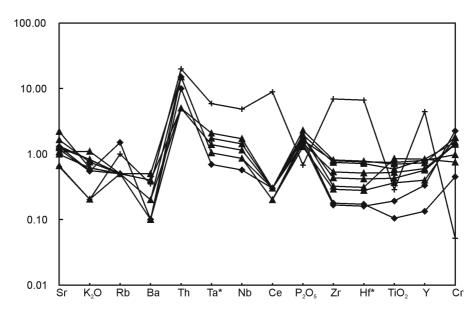


Figure 3. Spider-diagram normalized on MORB (Pearce 1983) for selected samples of the ophiolitic complex of the ŽOC. Explanation: Ta* and Hf* are calculated according to Rollinson (1993) - Ta*=Nb/16, and Hf*=Zr/39.

•: cummulitic rocks, •: gabbros and diabases, +: plagiogranite.

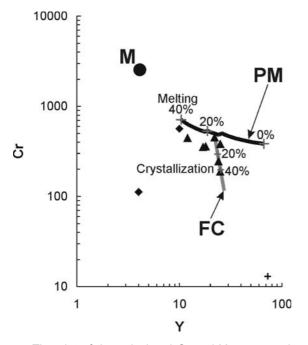


Figure 4. Partial melting model based on Cr and Y contents of the mantle source composition (M) and fractional crystallization model of a magma originated by partial melting. Line PM represents a non modal batch partial melting model within the range F=0.0 to 0.4 (0% - 40% melting). Line FC represents a fractional crystallization model within the range F=0.0 to 0.6 (0% - 40% crystallization). Symbols as in figure 3.

The plot of the calculated Cr and Y concentrations along with those of the analyzed samples supports that the Cr-richer samples could represent primary melts after 10 to 15% mantle-partial melting. Other samples show effects of modification processes.

The presence of glomeroporphyritic texture indicates that crystal fractionation was most important for magma modification. The major element model requires 40% (F=0.6) crystallization of a presumed primary magma (gabbro V-315) resulting in a mineral assemblage of $Pl_{52.9}Cpx_{12.5}Ol_{26.1}Ttn_{2.9}Ap_{4.4}Mgt_{1.0}$ to reproduce the major element composition of the more evolved

sample (diabase 8-A) (Tab. 2). Such a process explains satisfactorily the geochemical behaviour of most trace elements (Fig. 5).

Table 2. Major element modeling of the fractional crystallization process.

	- ,				,								
	Parental magma		Evolved magma		Composition of minerals crystallized								
-	obs calc			d ² =					-				
	Gabbro		Diabase	(obs-calc)2	PI	Срх	Fo	Fa	Tit	Ар	Mt		
Sample	V-315		8-A		V-315 E-536d		*	*	*	* E-536d			
SiO ₂	48.64	48.87	52.55	0.05	49.36	51.73	41.85	30.09	29.28	0.03	0.00	F	60%
TiO ₂	1.05	1.18	1.12	0.02	0.00	0.70	0.07	0.00	38.78	0.06	3.32		
Al_2O_3	15.73	15.59	14.69	0.02	31.59	1.35	0.00	0.00	1.67	0.02	0.53	PΙ	52.9
FeO _t	9.80	9.72	10.27	0.01	0.20	9.49	2.05	69.42	1.18	0.00	86.93	Срх	12.6
MnO	0.20	0.16	0.18	0.00	0.00	0.30	0.21	0.28	0.45	0.00	0.00	Fo	17.0
MgO	9.94	9.79	8.66	0.02	0.06	14.27	56.17	0.91	0.21	0.00	0.00	Fa	9.1
CaO	12.03	11.68	10.11	0.13	14.85	21.50	0.00	0.08	28.25	59.58	0.00	Tit	2.9
Na ₂ O	2.20	2.09	2.13	0.01	3.72	0.46	0.00	0.00	0.13	0.00	0.00	Аp	4.4
K ₂ O	0.12	0.06	0.09	0.00	0.03	0.00	0.00	0.00	0.05	0.20	0.00	Mt	1.0
P_2O_5	0.28	0.83	0.19	0.30	0.00	0.00	0.00	0.00	0.00	40.10	0.00		
$\Sigma d^2 = 0.56$													

^{*} from Deer et al. (1992).

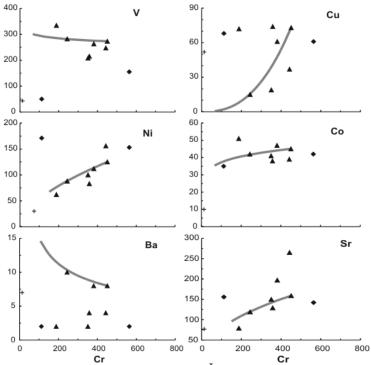


Figure 5. Diagrams showing trace element variations of the ŽOC basic rocks. Lines represent a fractional crystallization model within the range F=0.0 to 0.6 (0% - 40% crystallization) of an assemblage $Pl_{52.9}Cpx_{12.5}Ol_{26.1}Ttn_{2.9}Ap_{4.4}Mgt_{1.0}$ using partition coefficients from the literature. Symbols as in figure 3.

6.2 Geotectonic setting

The available analytical data at the moment cannot provide an unambiguous answer about geotectonic setting of the ŽOC. Analysed gabbro and diabase samples, which have been plotted on Nb*2-Zr/4-Y (Meshede 1986) and V vs. Ti (Shervais 1982) diagrams occupy both MORB and VAB fields. The same is confirmed by chemical compositions of clinopyroxenes plotted on the Ti+Cr vs. Ca (Letterier at al. 1982) and F_1 vs. F_2 (Nisbet & Pearce 1977). More precise data could be expected in future, after analyzing REE contents and isotope composition of the ŽOC rocks.

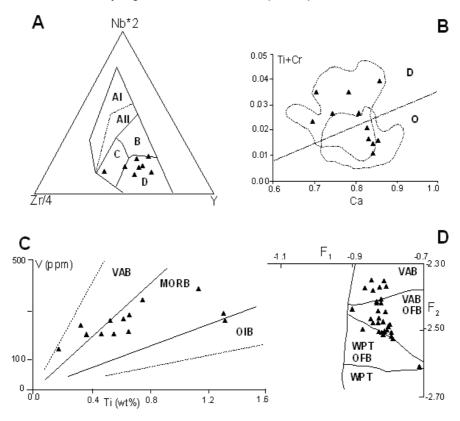


Figure 6. Diagrams for analyzed rocks (a-c) and clinopyroxenes (b-d) showing the geotectonic setting of the ŽOC. **A.** Nb*2-Zr/4-Y (Meshede 1986). Al: within-plate alkali basalts, All, within-plate alkali basalts and within-plate tholeiites, B: E-type MORB, C: within-plate tholeiites and volcanic-arc basalts, D: N- type MORB and volcanic-arc basalts. **B.** Ti+Cr vs. Ca (Letterier et al. 1982). D: field of MORB and other tholeiites from a spreading zone, O: volcanic arc basalts. **C.** V vs. Ti (Shervais 1982). VAB: volcanic arc basalts, MORB: mid-ocean ridge basalts, OIB: ocean island basalts. **D.** F₁ vs. F₂ (Nisbet & Pearce 1977). VAB: volcanic arc basalts, OFB: ocean floor basalts, WPT: within-plate tholeiites. **▲**: diabases and gabbros

7 SUMMARY

The ophiolitic complex of Ždraljica belongs to the Eastern Branch of the Vardar Zone Composite Terrane. It is composed mostly of gabbros and diabases with small occurrences of serpentinised wherlites, basalts and plagiogranites. This tholeiltic suite of rocks is intruded by intermediate and acid magmas of calc-alkaline character.

The emplacement of the ŽOC was before Upper Jurassic. Available K/Ar data gave a probable age of crystallization of about 168 Ma. The time of different metamorphic events are also recorded.

The tholeitic magmas of the ŽOC may have been formed by partial melting of the spinel-lherzolite (Ol₅₇Opx_{25.5}Cpx₁₅Sp_{2.5}) source. The results of a non-modal batch melting imply that the most primitive samples of the ŽOC have similar composition to melts formed by 10-15 % of partial

melting of such a source. More evolved magmas are modified by fractional crystallization (F=0.6) with the following fractionating assemblage: $Pl_{52.9}Cpx_{12.5}Ol_{26.1}Ttn_{2.9}Ap_{4.4}Mgt_{1.0}$.

Geochemical characteristics show that tholeiitic magmas have a MORB and VA affinity. Therefore, determination of a geotectonic setting cannot be done precisely.

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