Geochemistry of the Panjal Traps basalts (NW Himalaya): records of the Pangea Permian break-up

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Key words. - Panjal Traps, Continental flood basalts, Geochemistry, Neotethys, Rifting.

Abstract. - The late Lower to Middle Permian Panjal Traps (NW Himalaya, India-Pakistan) represent the greatest magmatic province erupted on the northern Indian platform during the Neotethys opening. New geochemical and isotopic analyses were performed on basalts from the eastern borders of the traps (SE Zanskar-NW Spiti area) in order to characterize this volcanism, to discuss its compositional variations in comparison to Panjal counterparts and its relationships with the opening of Neotethys. Lavas show features of tholeitic low-Ti (< 1.6%) continental flood basalts with LREE, Th enrichments and Nb-Ta negative anomalies. Trace element ratios combined with ENd, values (-3.6 to +0.9) and high Pb isotopic ratios suggest that these tholeitic basalts were derived from an OIB-like mantle contaminated at various degrees by a continental crust component. Previous geochemical features are broadly similar to those of the coeval Panjal volcanic sequences identified westwards (Ladakh, Kashmir and Pakistan). Present geochemical constraints obtained for the Panjal Traps basalts suggest they originated from rapid effusion of tholeitic melts during opening of the Neotethys Ocean. Similar magmatism implying an OIB-type reservoir is contemporaneously recognized on and along the adjacent Arabian platform. Both Indian and Arabian Permian volcanics were emplaced during coeval syn-rift to post rift transition. These Lower to Middle Permian south Neotethyan continental flood magmatism are regarded as associated to a passive rifting. In this scheme, OIB-type isotopic signature would be related either to a melting episode of syn-rift upwelling mantle plumes or to a melting of a regional abnormally hot and enriched mantle.

Géochimie des basaltes des trapps de Panjal (NW Himalaya): un enregistrement de la fragmentation de la Pangée au Permien

Mots-clés. - Trapps de Panjal, Basaltes continentaux, Géochimie, Néotéthys, Rifting.

Résumé. - Les trapps de Panjal (fin Permien inférieur - Permien supérieur, NW Himalaya) représentent la plus grande province magmatique mise en place sur la plate-forme nord indienne pendant l'ouverture de la Néotéthys. De nouvelles analyses géochimiques et isotopiques ont été réalisées sur les coulées basaltiques de la bordure Est des trapps, dans la région du SE Zanskar-NW Spiti. Le but est de caractériser ce volcanisme et de le comparer avec les équivalents connus des trapps de Panjal, afin de discuter des variations géochimiques existantes et de mieux comprendre ses relations avec l'ouverture de la Néotéthys. Les laves analysées ont des compositions de basaltes tholéitiques pauvres en Ti (< 1.6%), montrant un enrichissement en terres rares légères et en Th, ainsi que des anomalies négatives en Nb et Ta. Ces signatures géochimiques sont typiques de basaltes tholéitiques continentaux. Les rapports en éléments traces, les valeurs d'∈Nd_i (-3.6 to +0.9) et les rapports isotopiques élevés du Pb suggèrent que ces tholéiites dérivent de la fusion d'un manteau de type OIB, contaminé à des degrés variés par un composant enrichi de type croûte continentale. La signature géochimique des coulées étudiées est très similaire à celle des séquences volcaniques équivalentes qui sont reconnues plus à l'Ouest (Ladakh, Kashmir et Pakistan). Les compositions géochimiques de l'ensemble des basaltes de Panjal suggèrent qu'ils proviennent de l'effusion rapide de liquides tholéiitiques associés à l'ouverture de l'océan Néotéthys. Un magmatisme similaire et impliquant également un réservoir de type OIB caractérise à la même période la formation de la marge adjacente nord arabe. Le volcanisme des marges indienne et arabe s'épanche au cours de la transition des périodes syn-rift à post-rift. La mise en place de ce magmatisme continental sud néotéthysien au cours du Permien inférieur à moyen est interprétée comme associée à un rifting de type passif. Dans ce contexte, les signatures isotopiques de type OIB peuvent être reliées à la fusion de panaches mantelliques syn-rift, ou bien à la fusion d'un manteau régional enrichi et anormalement chaud.

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INTRODUCTION

Opening of the Neotethys Ocean produced the fragmentation of Pangea and resulted in the separation of numerous Peri-Gondwana continental blocks from the northeast of Gondwana. Southern margins of this ocean comprised the northern rim of African, Arabian, Indian and Australian plates [Stampfli et al., 1991; Dercourt et al., 1993]. These new continental plate boundaries evolved as passive margins during the Mesozoic. Since closure of the oceanic realm during the Cretaceous and Tertiary, major parts of the southern Neotethyan margins were inverted and incorporated within the Alpine chains. Today, preserved fragments of those margins carry traces of the different within-plate magmatic episodes, which accompanied their formation. The latter are dispersed along the Tethyan orogenic belt (fig. 1a), from Australia, Himalaya, Oman, to Zagros and eastern Mediterranean chains (Syria, Turkey, Cyprus, Maury et al., [2008]). This paper is focused on the NW Himalayan volcanics, which accompanied the break-up of the northern Indian plate rift during the Permian (fig. 1b).

Voluminous volcanic flows were emitted on the NW Indian rifted platform, during the late Lower to Middle Permian. They represent the greatest volume of melts emplaced during the Neotethyan opening and are called the Panjal Traps (fig. 1b). Whereas this name was used to describe the 2500 m-thick volcanic succession from the Kashmir mountains [Pareek, 1976], now it includes all the coeval flows found from Pakistan to the west, to Ladakh and Zanskar suite to the east [fig. 1c; Honegger et al., 1982; Papritz and Rey, 1989; Gaetani et al., 1990; Spencer et al., 1995]. The different Panjal lavas occurrences have been unevenly studied or their geochemical characteristics and often only in order to establish correlations between the various Permian Traps outcrops.

In the present work, we determined Sr-Nd-Pb isotopic data and major and trace element analyses on representative samples (sixteen lavas) of Panjal Traps basalts of the Zanskar-Spiti area. The main aims were to determine the geochemical affinities and source characteristics of this Permian volcanism. These data are compared with the coeval Panjal magmatic flows of the Pakistan, Kashmir and Ladakh and with worldwide continental flood basalts geochemical features. This paper yields the first complete geochemical study of the Lower to Middle Permian magmatic remnants from the NW Indian plate break-up.

GEOLOGICAL SETTING

Analyzed samples of volcanic rocks were sampled in the SE Zanskar-NW Spiti area (NW Himalaya, SE Ladakh, fig. 1c), which is the easternmost Panjal Traps flows outcrop. They belong to the Tethyan zone sedimentary belt of the higher Himalaya, which includes the Late Paleozoic to Mesozoic northern Indian platform sequence (fig. 1c). The latter corresponds to the northern Indian passive margin successions, preserved in thrust sheets or nappes. This Tethyan zone faces the northern Indus-Yarlung suture zone (IYSZ, fig. 1c) units, including the Indian margin slope and basin sequences, and the Tethyan oceanic units of the Spontang and Nidar ophiolites [Mascle, 1985].

The Carboniferous to early Permian sedimentary deposits of the Tethyan zone mainly consist of successive terrigenous detrital accumulations, separated by local unconformities, and associated with scarce magmatism mainly of alkaline affinity [Garzanti et al., 1992, 1994, 1996a,b; Vannay, 1993; Spring et al., 1993]. These sedimentary successions are considered as the products of rift-shoulder erosion taking place during differential uplift of the Indian margin rifted zones [Bassoulet et al., 1980; Vannay and Spring, 1993; Garzanti et al., 1999]. The late Paleozoic rifting phase and its associated magmatic remnants are documented in Pakistan [Pogue et al., 1992], Zanskar [Vannay and Spring, 1993; Spring et al., 1993] and in the central part of the chain [Central Nepal, Garzanti et al., 1992; 1994; Sciunnach and Garzanti, 1996]. More voluminous volcanic extrusions are identified in the eastern and central parts of the Himalaya (fig. 1b). They consist of the late Carboniferous-Early Permian Abor volcanics (~1500 m of basaltic to andesitic flows and tuffs, Arunachal Pradesh, NE Syntaxis) [Bhat, 1984], and the late Lower Permian 300 m-thick tholeiitic Nar-Tsum spilites and Bhote Kosi basalts (Sakmarian-Roadian), which lie on top of the late Palaeozoic syn-rift successions [South Tibet, Garzanti and Scuinnach, 1997; Garzanti et al., 1999]. These tholeitic affinity sequences erupted during a subsidence stage of the Indian margin and are interpreted as the remainders of the continental break-up and opening of the Neotethys [Garzanti et al., 19991.

The slightly younger Panjal Traps magmatic event [Artinskian-Kazanian, Nakazawa *et al.*, 1975; Gaetani *et al.*, 1990] led to the emplacement of the greatest magmatic province in the NW Indian margin rifted platform (fig. 1c). Remnants of the Panjal Traps magmatism are found in more than 10^5 km^2 from the easternmost Zanskar-Spiti to NE Pakistan in the west [Honegger *et al.*, 1982; Papritz and Rey, 1989].

In the Zanskar-Spiti-Lahul studied area (SE Ladakh), Panjal volcanic flows belong to the Kuling detrital terrigenous group of Garzanti *et al.* [1996a, b], which exhibits a major basal unconformity observed in Spiti [Fuchs, 1982] and Zanskar [Vannay, 1993; Garzanti, 1993; Garzanti *et al.*, 1996a]. In this area, Panjal continental flood basalts [Honegger *et al.*, 1982; Vannay and Spring, 1993] are regarded as traps, which filled a rifted valley recognized as the Zanskar-Spiti synclinorium [Fuchs, 1982; Gaetani and Garzanti, 1991; Garzanti *et al.*, 1996a].

Panjal Traps are also described throughout all Ladakh [fig. 1c, N Zanskar in Singh et al., 1976; Suru area in Honegger et al., 1982] and in the Kashmir basin [fig. 1c; Pareek, 1976; Bhat and Zainuddin, 1978, 1979] where they reach a 2500 m maximum thickness. Here they consist in a basal unit made of intermediate to acid pyroclastic rocks [Pareek, 1976] overlain by massive aphyric basaltic flows with tholeiitic to slightly alkaline affinities [Singh et al., 1976; Honegger et al., 1982; Gupta et al., 1983; Vannay and Spring, 1993]. The intermediate and acid differentiated rocks constitute a minor part of the total volume of the extruded rock [Pareek, 1976]. They were also recognized in the western syntaxis of NE Pakistan [fig. 1c; Papritz and Rey, 1989; Spencer et al., 1995]. In this zone, remnants of this magmatism have been identified within the lesser and higher Himalaya series [respectively, Panjal and Salkhala tectonic units, Papritz and Rey, 1989; and Upper Kaghan area, Spencer et al., 1995]. They are exposed as feeder

dykes crosscutting the crystalline basement and Lower Palaeozoic cover and as magmatic flows interlayered in the Upper Palaeozoic/Lower Mesozoic cover [Tethyan Zone,

Spencer *et al.*, 1995]. They are widespread (up to 2 km-thick) and metamorphosed under eclogites and amphibolites metamorphic facies conditions.

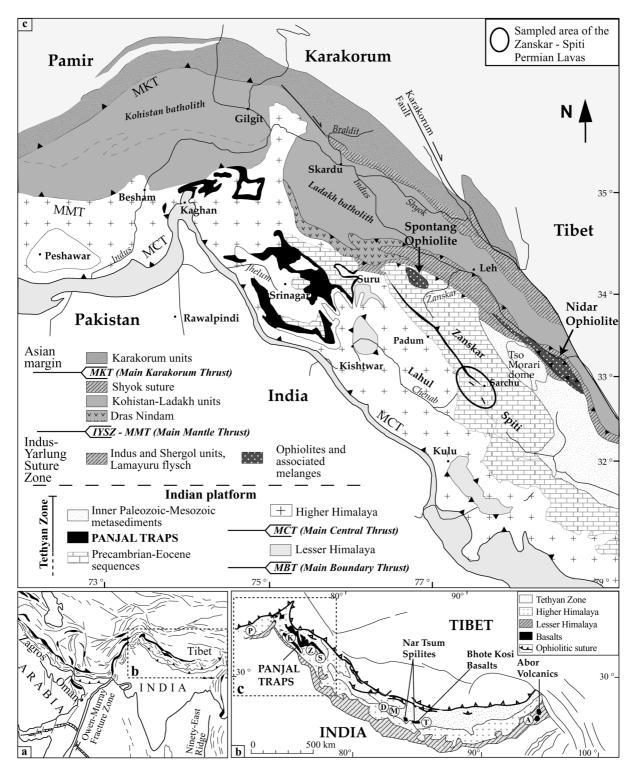


FIG. 1. – (a) Tethyan Suture (ophiolites and associated mélanges) after Coleman [1981]. (b) Main geologic features of the Himalayan Range showing the various occurrences of Permian basalts. P=North Pakistan; K=Kashmir; Z=Zanskar; S=Spiti; D=Dolpo; M=Manang, T=South Tibet; A=Arunachal Pradesh [Garzanti *et al.*, 1999]. (c) Geological map of the northwestern Himalaya modified after Guillot *et al.* [2003]; Spencer *et al.* [1995]; Pogue *et al.* [1992]; Vannay and Spring [1993].

FIG. 1. – (a) Suture téthysienne (ophiolites et mélanges associés) d'après Coleman [1981]. (b) Carte géologique simplifiée des principales unités de la chaîne himalayenne et montrant les différentes séquences basaltiques permiennes. (c) Carte géologique du NW de l'Himalaya, modifiée d'après Guillot et al. [2003]; Spencer et al. [1995]; Pogue et al. [1992]; Vannay et Spring [1993].

In the sampled area of the Zanskar-Spiti, traps eruption is followed by emplacement of the transgressive Kazanian to Djulfian sedimentary successions [upper succession of Kuling Group; Garzanti *et al.*, 1996a,b]. The latter testifies to the progressive thermo-tectonic subsidence of the Indian passive margin, which is interpreted as deriving form the Neotethyan oceanic expansion [Gaetani *et al.*, 1990; Gaetani and Garzanti, 1991; Garzanti *et al.*, 1999; Vannay, 1993].

The sixteen Panjal Traps basalts analyzed in the present work come from the sampling of Spring [1993] and Vannay [1993]. They selected samples in the Chumik and Zangla tectonic units located respectively in the SE Zanskar and

NW Spiti, in the northwestern Sarchu area, and along the Chandra Valley (High Lahul area), 30 km south of Sarchu [fig. 1c or see sampling map in Spring, 1993 and Vannay and Spring, 1993]. In these zones, the Panjal Traps are from 30 to 150 m thick and mainly crop out as massive flows, but also as pillow lavas and hyaloclastites [Spring, 1993].

ANALYTICAL PROCEDURES

Major elements were determined by Inductively Coupled Plasma-Atomic Emission Spectrometry (ICP-AES) at the Université de Bretagne Occidentale in Brest, following the

Table I. - Major and trace elements compositions for the Zanskar-Spiti Panjal Traps samples.

Tabl. I. - Compositions en éléments majeurs et en éléments traces des échantillons des trapps de Panjal du Zanskar-Spiti.

						- Grou	p 1 -								- Grou	ıp 2 -
Sample	Y96	Y99	Y100	Y101	Y102	Y103	Y104	Y105	Y117	Y118	Y130	04/08A	04/08B	793-13	293-11	293-12
SiO2 %	52.6	51.0	56.0	53.6	53.3	55.8	51.8	49.9	53.6	52.4	50.9	52.6	50.5	48.3	44.2	49.4
TiO2	1.23	1.44	1.53	1.28	1.21	1.59	1.28	1.33	1.19	1.19	1.17	1.20	1.32	0.71	0.52	0.62
Al2O3	15.6	17.5	15.0	15.6	15.3	16.6	15.8	16.7	15.9	15.6	15.3	15.7	17.3	16.1	13.7	14.6
Fe2O3	10.99	13.13	11.04	11.37	10.32	10.11	11.40	12.09	11.40	10.96	13.29	10.18	17.41	10.36	12.89	10.53
MnO	0.20	0.21	0.23	0.20	0.16	0.22	0.21	0.18	0.13	0.19	0.15	0.20	0.10	0.14	0.18	0.17
MgO	7.77	8.77	5.96	8.85	7.59	7.79	8.60	8.82	7.44	9.11	8.04	6.05	4.28	10.61	18.03	14.19
CaO	7.08	2.51	7.25	5.10	9.63	1.85	7.42	8.61	6.41	7.26	9.97	11.82	5.40	8.52	9.84	6.44
Na2O	4.04	5.23	2.88	3.81	2.36	5.83	3.36	2.27	3.42	3.15	1.05	1.61	2.59	4.03	0.48	3.55
K2O	0.46	0.07	0.03	0.03	0.02	0.05	0.03	0.03	0.42	0.03	0.01	0.53	1.01	1.06	0.03	0.39
P2O5	0.11	0.13	0.16	0.12	0.11	0.16	0.12	0.13	0.11	0.12	0.11	0.12	0.13	0.09	0.06	0.08
LOI	3.60	5.38	3.71	6.36	3.51	4.10	5.70	4.58	7.47	4.44	6.18	7.98	7.72	7.23	5.70	4.92
Sc (ppm)	40	42	43	37	37	46	38	40	36	36	34	36	38	47	36	44
Ti	7369	8633	9201	7659	7256	9540	7669	7985	7167	7142	7012	7189	7922	4288	3146	3748
v	278	300	310	270	270	325	335	285	265	262	244	272	260	191	230	170
Ni	92	102	57	96	90	76	95	95	89	92	86	85	135	195	920	305
Co	69	52	62	46	62	52	56	53	43	55	57	51	90	46	82	63
Cr	340	370	178	345	325	242	335	345	330	335	320	335	350	692	2440	905
Rb	16.0	1.0	0.3	0.4	0.5	0.8	1.0	0.6	13.3	1.2	0.6	16.8	41.1	44.0	0.4	20.0
Ba	186	81	29	41	43	38	41	37	388	54	34	252	198	254	18	152
Th	3.08	3.23	3.70	3.09	3.03	2.42	3.35	3.33	2.98	2.91	2.67	2.88	3.20	1.14	0.74	1.51
U	0.50	0.41	0.58	0.48	0.52	0.39	0.54	0.54	0.44	0.49	0.38	0.30	0.47	0.21	0.17	0.35
Nb	7.95	7.66	9.55	6.93	7.21	8.15	7.17	7.32	6.93	6.90	nd	6.78	6.95	2.35	1.63	2.60
Ta	0.83	0.57	1.15	0.57	1.06	0.70	0.63	0.68	0.54	0.62	nd	0.78	0.73	0.15	0.10	0.17
Pb	4.79	2.70	8.32	3.98	5.31	2.64	4.08	4.59	6.06	4.97	5.67	4.84	2.27	1.39	1.61	1.94
Sr	360.5	186.5	691.3	235.4	181.4	166.7	389.9	213.2	100.0	365.4	353.7	340.1	209.5	122.1	317.5	179.7
Zr	114.0	106.6	127.7	107.2	101.4	111.6	107.5	107.7	101.4	101.4	96.9	86.4	95.6	45.4	30.6	44.6
Hf	2.69	2.83	3.23	2.65	2.58	2.93	2.71	2.79	2.58	2.60	2.38	2.20	2.52	1.23	0.98	1.29
Y	25.69	23.30	26.48	23.60	22.64	25.26 8.19	24.03	23.88	22.64	23.06	19.98 10.53	23.83	20.02	25.02	18.74	26.87
La (ppm) Ce	12.25 26.52	10.68 25.23	14.01 30.93	8.70 22.11	11.46 24.89	21.77	10.80 24.10	12.59 27.29	12.07 26.15	10.81 24.20	23.83	11.92 25.47	9.94 21.09	4.18 8.73	2.25 5.37	4.72 10.62
Pr	3.42	3.25	4.01	3.05	3.24	3.11	3.22	3.51	3.55	3.24	3.08	3.28	3.01	1.21	0.74	1.43
Nd	14.53	14.05	16.78	13.66	13.70	14.04	13.79	14.72	14.89	13.80	13.13	13.70	13.12	5.64	3.99	6.71
Sm	3.47	3.56	4.02	3.35	3.29	3.87	3.57	3.62	3.58	3.37	3.14	3.51	3.43	1.98	1.49	2.48
Eu	1.08	0.96	1.35	1.06	1.08	1.09	0.99	1.19	1.17	1.06	0.99	1.10	0.97	0.79	0.63	0.77
Gd Tb	3.94 0.65	3.99 0.68	4.34 0.74	3.99 0.65	3.62 0.61	4.17 0.70	3.93 0.64	4.19 0.68	4.20 0.69	3.69 0.62	3.29 0.54	3.87 0.63	3.63 0.61	2.77 0.53	2.15 0.42	3.06 0.64
Dy	3.99	3.87	4.48	3.86	3.72	4.22	4.12	4.02	4.24	3.70	3.27	3.75	3.67	3.65	3.15	4.42
Ho	0.80	0.81	0.93	0.84	0.79	0.90	0.86	0.87	0.88	0.81	0.68	0.79	0.74	0.84	0.72	1.00
Er	2.30	2.34	2.68	2.23	2.17	2.54	2.51	2.46	2.41	2.15	1.87	2.15	2.13	2.51	2.08	2.91
Yb	2.12	2.09	2.52	2.08	1.98	2.33	2.24	2.29	2.09	1.94	1.61	1.84	1.90	2.43	2.05	2.83
Lu	0.31	0.33	0.37	0.31	0.31	0.35	0.34	0.34	0.32	0.29	0.25	0.27	0.27	0.37	0.32	0.42
La/Nb (La/Yb) _N	1.54 4.14	1.39 3.67	1.47 3.99	1.26 3.00	1.59 4.15	1.00 2.52	1.51 3.46	1.72 3.94	1.74 4.14	1.57 4.00	nd 4.69	1.76 4.65	1.43 3.75	1.78 1.24	1.38 0.79	1.82 1.20
(La/Tb) _N (La/Gd) _N	2.70	2.32	2.80	1.89	2.74	1.70	2.38	2.61	2.49	2.54	2.78	2.67	2.38	1.24	0.79	1.34
Eu/Eu*	0.89	0.78	0.99	0.89	0.96	0.83	0.81	0.93	0.92	0.92	0.94	0.91	0.84	1.03	1.08	0.85
Nb/Y	0.31	0.33	0.36	0.29	0.32	0.32	0.30	0.31	0.31	0.30	nd	0.28	0.35	0.09	0.09	0.10
Th/Ta	3.71	5.67	3.22	5.42	2.86	3.46	5.32	4.90	5.52	4.69	nd	4.26	4.36	7.72	7.40	8.88
Mg#	0.58	0.57	0.52	0.61	0.59	0.60	0.60	0.59	0.56	0.62	0.54	0.54	0.33	0.67	0.73	0.73

Major elements, Sc, V, Ni and Cr contents were determined by ICP-AES and others trace elements by ICP-MS Major element contents are recalculated volatile free $Eu/Eu^* = EuN / [(SmN^*GdN)]1/2$

procedures of Cotten *et al.* [1995] and using international standards for calibration tests (ACE, BEN, JB-2, PM-S, WS-E). Rb was measured by flame atomic emission spectroscopy. Relative standard deviations are ca. 1% for SiO_2 and 2% for other major elements except P_2O_5 and MnO (0.01%). All rocks were pulverised in an agate mill.

Trace elements including the REE were analyzed by ICP-MS at the Université Joseph Fourier in Grenoble, after acid dissolution of 100 mg sample, using the procedures of Barrat *et al.* [1996]. Standards used for the analyses were BHVO and BIR-1. Analytical errors are less than 5% for trace elements.

Sr (static acquisition) and Nd (semi-dynamic acquisition) isotopic ratios were measured at the LMTG de l'Université Paul Sabatier de Toulouse on a Finnigan MAT 261 multicollector mass spectrometer using the analytical procedures of Lapierre et~al.~[1997]. Results on standards yielded $^{143}\mathrm{Nd}/^{144}\mathrm{Nd}=0.511961\pm9~(n=4)$ and $^{87}\mathrm{Sr}/^{86}\mathrm{Sr}=0.710252\pm9~(n=4)$ for neodynium rennes standard [Chauvel and Blichert-Toft, 2001] and NBS 987, respectively. $^{143}\mathrm{Nd}/^{144}\mathrm{Nd}$ and $^{87}\mathrm{Sr}/^{86}\mathrm{Sr}$ were normalized for mass fractionation relative to $^{146}\mathrm{Nd}/^{144}\mathrm{Nd}=0.7219$ and $^{88}\mathrm{Sr}/^{86}\mathrm{Sr}=8.3752,$ respectively. Two leaching steps (HCl acid during 20 minutes at $100^{\circ}\mathrm{C})$ were done before acid digestion of samples.

Nine basalts were selected for lead separation. Selection was conducted after trace element measurements and neodynium isotopic analyses, in order to choose the greatest chemical and isotopic diversity of the sample set. A leaching

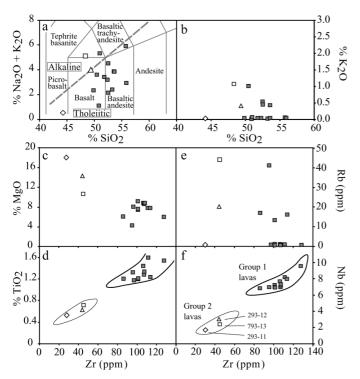


FIG. 2. — (a) Na₂O + K₂O (%) vs SiO₂ (%) [MacDonald and Katsura, 1964], (b) K₂O (%) vs Zr (ppm), (c) Mg0 (%) vs Zr (ppm), (d) TiO₂ (%) vs Zr (ppm), (e) Rb (ppm) vs Zr (ppm), (f) Nb (ppm) vs Zr (ppm) plots for the Zankar-Spiti samples of the Panjal Traps. Legend is reported in the fig. 2f. FIG. 2. — Diagrammes de correlation : (a) Na₂O + K₂O (%) vs SiO₂ (%) [MacDonald et Katsura, 1964], (b) K₂O (%) vs Zr (ppm), (c) Mg0 (%) vs Zr (ppm), (d) TiO₂ (%) vs Zr (ppm), (e) Rb (ppm) vs Zr (ppm), (f) Nb (ppm) vs Zr (ppm) pour les échantillons des trapps de Panjal du Zanskar-Spiti. La légende est indiquée sur la fig. 2f.

step with 6N HCl during 30 minutes at 65°C was done before acid digestion (36-48 hours on a hotplate in ultrapure HF and HNO₃ concentrate acids). After complete evaporation, 0.5 ml of concentrated HBr was added to the sample and kept at 70°C for 2-3 hours before complete evaporation. The chemical separation of lead was done using 50 µl of anion exchange resin (AG1X8, 200-400 mesh) with 0.5N HBr and HCl acids as reagents. Pb blanks were less than 40 pg. Lead isotope measurements were made on a VG model Plasma 54 multi-collection magnetic sector-inductively coupled plasma-mass spectrometer (MC-ICP-MS) at the Ecole Normale Supérieure in Lyon. Lead isotope compositions were measured using the Tl normalization method described by White et al. [2000]. For Pb isotope analyses, samples were bracketed between NBS 981 standards and calculated with respect to the value reported for this standard by Todt et al. [1996]. This technique yields internal precision of ca. 50 ppm (2s) and an external reproducibility of ca. 250 ppm (2s) for 206 Pb/ 204 Pb ratios (n=15). Initial isotopic ratios have been calculated assuming an age of 275 Ma.

RESULTS

Petrography

Most of the samples (collected by Spring and Vannay in 1993) were already powdered before being analyzed. In the three available rock samples, magmatic minerals have been modified by the Himalayan metamorphism and weathering processes. Two out of the sixteen Panjal basalts (793-13 and 293-12), have preserved aphyric, spherulitic and vesicular textures of pillow rims. The plagioclase microlites organized in spherulites are albitized. The groundmass is mainly constituted by the following assemblage albite + oxydes + chlorite + epidote + actinolite + calcite, which characterizes a greenschist facies previously recognized by Vannay and Spring [1993]. Sample 293-11 shows a porphyric texture with numerous pseudomorphs of olivine phenocrysts, replaced by serpentine and surrounded by an assemblage made of chlorite, epidote and actinolite. The latter likely derives from destabilisation of clinopyroxene. The high Cr (2440 ppm) and Ni (920 ppm) contents of the whole rock suggest initial accumulation of olivine and clinopyroxene (table I).

Major and trace elements

The loss of ignition (LOI) of analysed samples ranges between 3.5 and 8% (table I). These high contents suggest significant modification of the Zanskar-Spiti Permian lavas primary composition. This is also evidenced with the erratic behaviours that are observed plotting the mobile major elements Na₂0, CaO, and K₂0 against SiO₂ contents variations (figs. 2a-b). In the MacDonald and Katsura [1964] discriminating diagram (fig. 2a), analyzed lavas mainly fall in the tholeiites basic field, between the picritic SiO₂ content of the 293-11 cumulate and the basaltic andesite compositions for the most evolved melts. In this diagram, the scattering of data suggests that K₂O and Na₂O contents have been remobilized during secondary processes. The zirconium content has therefore been reported against major (figs. 2c-d), lithophile (fig. 2e) and incompatible high field strength elements (fig. 2f) in view to check the mobility of elements during

secondary processes. TiO_2 and Nb versus Zr contents exhibit good correlations and thus are considered as representative of initial magmatic characters. In contrast, no satisfactory alignment is obtained in Na_2O , K_2O (not shown), MgO, Sr and Rb versus Zr diagrams (figs. 2b-e) suggesting significant mobility of these elements during secondary processes and that they can not be used to characterize the magmatic affinities of the studied Panjal Traps layas.

Analyzed basalts can be subdivided into two main groups based on their ${\rm TiO_2}$, ${\rm Zr}$, Nb and REE contents (figs. 2-3). Group 1, which encompasses the large majority of Zanskar-Spiti samples (13 samples), is characterized by homogeneous Zr contents (86< ${\rm Zr_{ppm}}$ <128 with an average of 105 ± 20 ppm), significantly higher than those measured for Group 2 samples (793-13, 293-12 and 293-11; 31< ${\rm Zr_{ppm}}$ <45) (table I). The Panjal volcanics show basaltic contents in MgO (MgO>6%, excepted for sample 04/08 B) with low ${\rm TiO_2}$ (<1.6%) and ${\rm P_2O_5}$ (0.06< ${\rm P_2O_5}$ %<0.16) contents. Overall, Group 2 rocks have, compared to Group 1, significantly lower ${\rm SiO_2}$, ${\rm TiO_2}$ and ${\rm P_2O_5}$ contents and higher concentrations in MgO (10.6 to 18%, table I). Thus, Group 2 presents geochemical affinities with more primitive liquids than Group 1.

Group 1 chondrite-normalized REE patterns show homogeneous distributions with significant LREE enrichment relative to MREE (1.7<(La/Gd)_N<2.8) and HREE (2.5<(La/Yb)_N<4.7) and slight Eu negative anomalies (0.78<Eu/Eu*<0.99) (fig. 3a, table I). Group 1 primitive mantle-normalized multi-elements spectra are characterized by significant Th and LREE enrichment and by marked Nb negative anomalies (1.0<La/Nb<1.8) (fig. 3b, tables Ia-b). No significant depletion of the less incompatible elements is observed. Samples 793-13 and 293-12 (Group 2) show rather flat chondrite-normalized REE-patterns ((La/Yb)_N ~ 1.2) with small LREE enrichment while sample 293-11 has a slight LREE depleted pattern ((La/Yb)_N = 0.8). Group 2 is

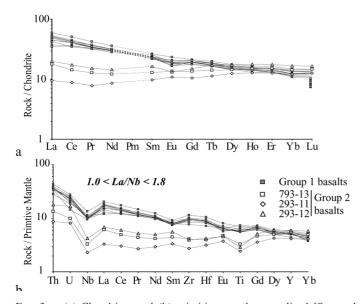


FIG. 3. – (a) Chondrite– and (b) primitive mantle-normalized [Sun and McDonough, 1989] patterns of the Panjal Traps basalts.
FIG. 3. – Composition en éléments traces des basaltes de Panjal: (a) Spectres de terres rares normalisés aux chondrites et (b) spectres multi-éléments normalisés au manteau primitif [d'après Sun et McDonough, 1989].

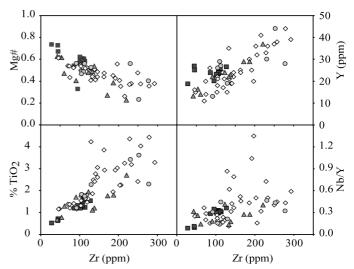
also enriched in Th but displays highest negative anomalies in Nb (La/Nb ~ 1.8) and Ti.

Comparison with the western Panjal Traps counterparts

Pioneer major element geochemical analyses from Kashmir and Zanskar basalts have shown that Panjal Traps flows have mainly tholeiitic affinities [Pareek, 1976; Singh *et al.*, 1976]. Major and trace elements works on coeval Permian volcanic rocks (basaltic flows and feeder intrusive bodies) outcropping in Pakistan and Ladakh have been plotted together with our data in view to establish geochemical composition variations through the Panjal Traps lavas from the Zanskar-Spiti to the Pakistan (figs. 4-5a)

Mg# [Mg/(Mg+Fe²⁺)], TiO₂ (wt%) and Y (ppm) show relatively good correlations with Zr (fig. 4). This indicates that the variation in element contents of the Panjal volcanics is more probably related to igneous fractionation processes rather than to metamorphic events. Group 1 Mg# values and Zr contents indicate basaltic compositions (51.2<Mg#<62.1), excepted for basalt 04/08B (Mg#=32.6) showing no SiO₂ content correlation (table I). Group 2 exhibits significantly higher Mg# values (66.8<Mg#<73.4) correlated with lower SiO₂ amount, this is assigned to a more primitive character of these lavas. The highly magnesian 293-11 sample, (MgO = 18%, Ni = 850 ppm, Cr = 2440 ppm), should represent the remnant of an ultramafic cumulate (olivine, clinopyroxene).

Most of the samples collected in Kashmir, Zanskar and Spiti show low TiO₂, Zr (fig. 4), and P₂O₅ (fig. 5a) contents. The greatest chemical variability is observed in the western syntaxis lavas (NE Pakistan) which display features of tholeitic to slightly alkaline affinities, characterized by larger scale Zr and TiO₂ variations for broadly similar Mg# [Pogue *et al.*, 1992; Spencer *et al.*, 1995]. Despite a



- SE Zanskar-NW Spiti [this paper]

 NE Pakistan [Spencer et al., 1995]
- NW Ladakh [Honegger et al., 1982; Spencer et al., 1995]

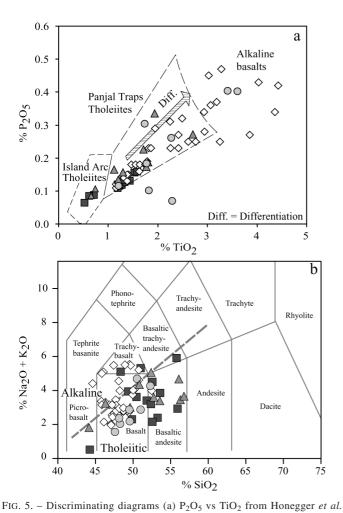
 NE Pakistan [Papritz and Rey, 1989]
- FIG. 4. Geochemical composition of the Panjal Traps lavas plots in Mg#, ${\rm TiO_2},~{\rm Y}$ and Nb/Y vs Zr diagrams.

FIG. 4. – Composition géochimique des laves des trapps de Panjal représentée dans les diagrammes de corrélation Mg#, TiO₂, Y et Nb/Y en fonction de Zr.

probable erratic behaviour of alkali major elements during secondary processes (fig. 5b), the discriminating diagram of MacDonald and Katsura [1964] also illustrates that most of the available Panjal lavas analyses are characterized by mainly basaltic compositions which plot within the tholeiitic melts field (fig. 5b).

Isotopic compositions of Zanskar-Spiti tholeiites

Zanskar-Spiti tholeiites (Groups 1 and 2) show very high and anomalous initial Sr isotopic ratios ranging from 0.7083 to 0.7144 (tables IIa-b). Such high and anomalous values can be assigned to an insufficient "in situ" decay correction due to mobility of lithophile elements during late alteration processes and especially the leaching of Rb. The extremely low Rb contents shown by some of the studied lavas (Rb_{ppm} < 1, fig. 2e, tables IIa-b) substantiate this hypothesis. εNd_i values range between +0.9 and -3.6 with six out of nine samples which have negative values. Such isotopic



[1982] and (b) [Na₂O+K̄₂O] vs SiO₂ diagram from MacDonald and Katsura [1964] for the Panjal Traps lavas. See legend in figure 4. In the figure 5a, Panjal tholeiitic field is from Honegger et al. [1982]; alkaline basalts and island arc tholeiites fields are from Hawkins [1980]. In the figure 5b, the stippled line distinguishes tholeiitic from alkaline basalts. FIG. 5. – Diagrammes discriminants (a) P₂O₅ vs TiO₂ d'après Honegger et al. [1982] et (b) [Na₂O+K₂O] vs SiO₂ d'après MacDonald et Katsura [1964] pour les laves des trapps de Panjal. Légende identique à la figure 4. Sur la figure 5a, le champ des tholéiites de Panjal est défini par Honegger et al. [1982]; et les champs discriminants des basaltes alcalins et des tholéiites d'arc insulaire sont d'après Hawkins [1980]. Sur la figure 5b, la

ligne pointillée distingue les basaltes tholéitiques des basaltes alcalins.

signatures ruled out the depleted mantle (DMM) as the major component involved in the source of the studied volcanic rocks.

In the Pb-Pb correlation diagrams (figs. 6a-b), Zanskar-Spiti tholeiitic basalts plot significantly above the North Hemisphere reference line [Zindler and Hart, 1986]. For a given (206Pb/204Pb) ratio, the two Zanskar-Spiti tholeiitic $(^{207}\text{Pb/}^{204}\text{Pb})$ show more radiogenic groups (208Pb/204Pb) values than both Indian and Atlantic MORBs (figs. 6a-b). In the (²⁰⁸Pb/²⁰⁴Pb)_i vs (²⁰⁶Pb/²⁰⁴Pb)_i plot, tholeiites report in the OIB field whereas in the (207Pb/204Pb), versus (206Pb/204Pb)_i diagram (fig. 6a), a significant shift of the data is observed towards a ²⁰⁷Pb-rich component close to the Enriched Mantle II-type component [EM II; Zindler and Hart, 1986]. Reported in the ENd_i-(206Pb/204Pb)_i diagram (fig. 6c), Group 1 and 2 basalts plot again between the OIB-type mantle and EM II-type enriched component (table II) with the exception of sample 293-11. The Group 2 basalts show higher initial Pb isotopic ratios than Group 1 (tables IIa-b). Samples from Group 1 show very homogeneous isotopic compositions for both the Nd and Pb isotopes.

DISCUSSION

Source origin for the Zanskar-Spiti Permian basalts

Pb and Nd isotopic ratios show that the studied easternmost Panjal Traps flows derived from the melting of an OIB-like mantle source. High Pb initial ratios and negative ENd; values also strongly suggest involvement of an enriched component. Plotted in the ¹⁴⁷Sm/¹⁴⁴Nd ratios versus εNd; diagram, the two groups of Panjal Traps tholeiites define two separated fields with significantly higher ¹⁴⁷Sm/¹⁴⁴Nd ratios for group 2 (fig. 7a). These latter basalts are probably generated by a higher degree of partial melting than group 1 and from a mantle reservoir defined by an ENd; value close to +1 (i.e. higher εNd_i values for the group 2 basalts). This assumption is consistent with the lower (La/Yb)_N and (La/Gd)_N ratios shown by group 2 compared to group 1 (tables I). Moreover, the isotopic compositions of the two groups of samples suggest that some contamination process, at variable degree, occurred between an OIB-type mantle source and an enriched reservoir (i.e. enriched lithospheric mantle, continental crust, terrigenous recycled sediments...). This contamination induces an increase of the Pb/Pb isotopic ratios simultaneously with a decrease of 143Nd/144Nd values (fig. 6c). Such an evolution is consistent with participation of an EM II-type contaminant, which is characterized in particular by high ²⁰⁷Pb/²⁰⁴Pb and moderate 206Pb/204Pb ratios and relatively low Nd isotopic ratios (fig. 6).

Plotted in the εNd_i versus Zr content diagram (fig. 7b), again the two previous groups are individualized and group 1 provides a more homogeneous domain than group 2. Both groups are characterized by relatively homogeneous Zr contents (table I). Two out of the three samples belonging to group 2 yield the lowest Zr contents coupled to the lowest εNd_i values of the present study. Therefore, the group 1 scattered field does not characterize any correlation between εNd_i and Zr contents. Thus, the contribution of an enriched component in the genesis of the studied lavas probably did not occur during fractional crystallisation into

TABLE IIa. – Nd, Sr and Pb isotopic compositions of the Zanskar-Spiti Panjal Traps samples. TABL. IIa. – Compositions isotopiques du Nd, du Sr et du Pb, des échantillons des trapps de Panjal du Zanskar-Spiti.

			- Group 1 -			
Sample	Y101	Y103	Y104	Y105	Y117	Y118
Db ()	0.25	0.79	0.05	0.50	12.24	1.15
Rb (ppm)	0.35	0.78	0.95	0.59	13.34	1.15
Sr (ppm)	235.41	166.73	389.88	213.21	99.96	365.36
⁸⁷ Sr/ ⁸⁶ Sr	0.714037 ± 10	0.712762 ± 10	0.710771 ± 9	0.710907 ± 7	0.715115 ± 7	0.714455 ± 9
⁸⁷ Rb/ ⁸⁶ Sr	0.004	0.014	0.007	0.008	0.386	0.009
$(^{87}Sr/^{86}Sr)_{i}$	0.71402	0.71271	0.71074	0.71088	0.71360	0.71442
ESri	139.8	121.2	93.3	95.1	133.9	145.5
Sm (ppm)	3.35	3.87	3.57	3.62	3.58	3.37
Nd (ppm)	13.66	14.04	13.79	14.72	14.89	13.80
¹⁴³ Nd/ ¹⁴⁴ Nd	0.512504 ± 5	0.512628 ± 5	0.512588 ± 4	0.512512 ± 5	0.512517 ± 6	0.512496 ± 5
¹⁴⁷ Sm/ ¹⁴⁴ Nd	0.148	0.167	0.157	0.149	0.145	0.148
$(^{143}Nd/^{144}Nd)_{i}$	0.51224	0.51233	0.51231	0.51224	0.51226	0.51223
ENd	-0.91	0.86	0.43	-0.77	-0.56	-1.05
²⁰⁶ Pb/ ²⁰⁴ Pb	18.530	18.509	18.497	18.486	18.433	18.457
207 Pb/ 204 Pb	15.690	15.684	15.684	15.672	15.673	15.672
208 Pb/ 204 Pb	39.324	39.148	39.166	39.121	39.119	39.093
$(^{206}Pb/^{204}Pb)_{i}$	18.20	18.10	18.13	18.16	18.23	18.19
$(^{207}\text{Pb}/^{204}\text{Pb})_{i}$	15.67	15.66	15.66	15.65	15.66	15.66
$(^{208}\text{Pb}/^{204}\text{Pb})_{i}$	38.62	38.32	38.42	38.47	38.68	38.57

shallow level magma chambers. This process can occur either in the early stages of magma evolution (i.e. mantle source) or later, during magma ascent to the surface.

Melts derived from primitive mantle are expected to have a Th/Ta ratio of ~ 2.1 [Sun and McDonough, 1989]. In the Th versus Ta diagram (fig. 7c), the group 1 and group 2 defined for the Panjal basalts plot along two distinct correlation lines defined by Th/Ta ratios of ~ 5.7 and ~ 8.2 , respectively. All basalts from group 2 are located on the Th/Ta line equal to ~ 8.2 suggesting that they derive from a same parental magma. Group 1 basalts show a rather large range of Ta values coupled with homogeneous Th abundance. However, three samples (Y100, Y102 and Y103, fig. 7c, table I) are more homogeneous and are located on the single Th/Ta = 3.2 correlation line. Such behaviour indicates the absence of any significant fractionation process and should reflect variable contamination degrees by an enriched

TABLE IIb. – Nd, Sr and Pb isotopic compositions of the Zanskar-Spiti Panjal Traps samples.

Tabl. IIb. – Compositions isotopiques du Nd, du Sr et du Pb, des échantillors des trapps de Panjal du Zanskar-Spiti.

		- Group 2 -		
Sample	793 - 13	293-11	293-11dup.	293-12
Rb (ppm)	43.96	0.40	0.40	19.97
Sr (ppm)	122.15	317.47	317.47	179.74
87Sr/86Sr	0.712374 ± 8	0.712149 ± 10	0.712116 ± 8	0.712197 ±
87Rb/86Sr	1.042	0.004	0.004	0.322
$(^{87}Sr/^{86}Sr)_{i}$	0.70830	0.71213	0.71210	0.71094
εSr_i	58.5	113.0	112.5	96.0
Sm (ppm)	1.98	1.49		2.48
Nd (ppm)	5.64	3.99		6.71
143Nd/144Nd	0.512588 ± 6	0.512722 ± 9		0.512504 ± 1
147Sm/144Nd	0.212	0.226		0.223
(143Nd/144Nd);	0.51221	0.51232		0.51210
ENd	-1.51	0.62		-3.56
206 Pb/ 204 Pb	19.000	19.086		19.023
207 Pb/ 204 Pb	15.784	15.767		15.745
208 Pb/ 204 Pb	39.563	39.438		39.441
$(^{206}\text{Pb}/^{204}\text{Pb})_{i}$	18.58	18.79		18.52
$(^{207}\text{Pb}/^{204}\text{Pb})_{i}$	15.76	15.75		15.72
$(^{208}\text{Pb}/^{204}\text{Pb})_{i}$	38.81	39.02		38.73

component. This is also suggested by the broadly reverse correlation existing between ϵNd_i values and La/Nb ratios (fig. 7d). The latter indicates a relationship between the extent of the Nb negative anomaly and the amount of the enriched component added to the OIB-like mantle. Moreover, the average Th/Ta value of the upper continental crust is as high as 10.7 [McLennan, 2001]. Taking into account isotopic and trace elements features of the studied basalts, it should be envisaged that parental magmas of both Groups 1 and 2 have been contaminated at some stages of their evolution by upper continental crust.

Spencer et al. [1995] published a set of Sr and Nd isotopic analyses from Pakistan Panjal Traps. The corresponding isotopic field is plotted against the Ti/Y ratio (fig. 8). In the absence of Fe-Ti oxide fractionation, the Ti/Y ratio is not affected by low-pressure fractionation processes [Brewer et al., 1992] and so, variations of this ratio can be attributed either to a variable degree of partial melting or to a mixing between various components. The latter process will be characterized by significantly different Ti/Y values. In the εNd_i vs Ti/Y diagram (fig. 8), Panjal Traps lavas of Zanskar-Spiti and Pakistan plot between a crust-derived end-member and an OIB-like mantle component. The broad tendency of our data (excepted for the group 2) suggests that the mantle component initially involved in the Panjal basalts was characterized by higher Ti/Y values than normal melts generate in mid-oceanic ridge. The scattering of data can be tentatively related to drastic change in melting conditions, especially the group 2 Panjal basalts characterized by lower Ti/Y ratios are related to a higher degree of partial melting. Moreover, the lowest ENdi values occur in the least metamorphosed and fractionated rocks of the NW Zanskar and Zanskar-Spiti Panjal tholeiites [Spencer et al., 1995]. This demonstrates that the crustal contamination process is more efficient in the more primitive high-temperature tholeiites (with exception of the 293-11 cumulate).

Crustal contamination processes have been often mentioned for the more primitive, higher temperature, magmas of continental flood basalts [Saunders *et al.*, 1992]. This is the case, for example, of the Poladpur and Bushe

formations from the Deccan Traps, which are characterized by high LILE (K, Rb, Ba...), Th and Pb-Sr isotopic ratios, and low Zr, Nb, Y contents and Nd isotopic ratios [Lightfoot et al., 1990; Lightfoot and Hawkesworth, 1988]. Similar features have been described for low-Ti basalts from various provinces such as the East Coast of Madagascar [Saunders et al., 1992], the Paraná province [i.e. Gramado basalts; Peate and Hawkesworth, 1996], the Guinean tholeiites [i.e. tholeiites from the Central Atlantic Magmatic Province; Deckart et al., 2005], and the Siberian traps basalts [i.e. Wooden et al., 1993; Sharma, 1997]. Alternatively, very low to intermediate ²⁰⁶Pb/²⁰⁴Pb values are often attributed to contribution of an enriched sub-continental lithospheric mantle component in the genesis of the basalts [e.g. Saunders et al., 1992; Kieffer et al., 2004].

The differences in the Nd and Pb isotopic compositions of the studied Panjal Traps basalts are not clearly linked to AFC process (fig. 7a) but more probably reflect contamination of an OIB-like mantle by an enriched component (figs. 7a-c-d). Some arguments allow us to propose that the

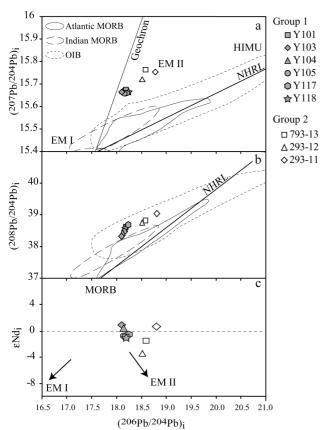


FIG. 6. – Isotopic compositions of the Panjal Traps basalts: (a) (207Pb/204Pb)_i vs (206Pb/204Pb)_i diagram. (b) (208Pb/204Pb)_i vs (206Pb/204Pb)_i diagram. (c) εNd_i vs (206Pb/204Pb)_i diagram. Fields of MORBs are from pre-compiled isotopic compositions in the PETDB (spreading axis tholeiitic basalts); Oceanic Islands Basalts (OIB) field, mantle components HIMU, Enriched Mantle EM-I and -II and the North Hemisphere Reference Line (NHRL) are from Zindler and Hart [1986].

FIG. 6. – Compositions isotopiques des basaltes des trapps de Panjal: Diagrammes (a) (207Pb/204Pb)_i vs (206Pb/204Pb)_i, (b) (208Pb/204Pb)_i vs (206Pb/204Pb)_i, (b) (208Pb/204Pb)_i vs (206Pb/204Pb)_i. Les champs de composition des MORBs correspondent aux compositions isotopiques pré-compilées des basaltes tholeiitiques échantillonnés à l'axe d'accrétion, sur le site de PETDB. Le champ de composition des basaltes des îles océaniques (OIB), des composantes mantelliques HIMU, EM-I et EM-II et la NHRL sont d'après Zindler et Hart [1986].

enriched component is the upper continental crust. They are (i) high Th/Ta and La/Nb ratios, (ii) negative correlation between ϵNd_i and Th/Ta ratios, and (iii) the geological context of Panjal Traps basalts emplacement, which is favourable to this process with an emplacement of these volcanics into the Indian continental crust.

Geodynamic implications

The chemical compositions of the Panjal Traps lavas suggest that effusions were largely dominated by basaltic melts of tholeitic affinity (figs. 5a-b). Their low TiO₂, Zr, and P₂O₅ contents and Nb/Y ratios (mostly lower than 0.6) but also the Th and LREE enrichments combined with Nb negative anomalies are typical features of continental flood tholeiitic basalts [Boillot and Coulon, 1998]. The minor proportion of high-fractionated melts [Pareek, 1976; Honegger et al., 1982; Papritz and Rey, 1989] and absence of Eu anomalies in Panjal lavas suggest a rapid effusion of magmas. This is commonly observed in many continental flood volcanic provinces where there is a striking contrast between the compositions of tholeiitic basalts and the petrographic and chemical diversity of other volcanic types [Arndt et al., 1998; Wilson, 1989; Iacumin et al., 2003]. This feature combined with the mainly tholeitic affinities observed in the wide geographic extent of the traps

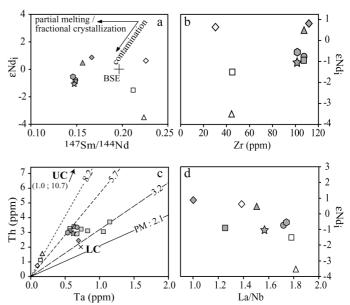


FIG. 7. – (a) ϵ Nd_i vs (147 Sm/ 144 Nd) diagram. (b) ϵ Nd_i vs Zr (ppm) diagram. (c) Th (ppm) vs Ta (ppm) diagram. Primitive mantle (PM) Th/Ta ratio (2.1) after Sun and McDonough [1989]; upper and lower continental crust (UC and LC) average contents after McLennan [2001]. Th/Ta ratio of each correlation line are plotted. (d) ϵ Nd_i vs (La/Nb) diagram. Samples are the same as those of the figure 6 and additional light grey squares in figure 7c correspond to the other analysed Zanskar-Spiti Panjal tholeites.

FIG. 7. – (a) Diagramme de corrélation εNd_i en fonction de $(1^{47}Sm/^{144}Nd)$. (b) Diagramme corrélation εNd_i en fonction de Zr (ppm). (c) Diagramme de corrélation des teneurs en Th (ppm) en fonction des teneurs en Th (ppm) pour les tholéittes de Panjal. Le rapport Th/Ta (2.1) du manteau primitif (PM) est d'après Sun et McDonough [1989]; les compositions moyennes de la croûte continentale supérieure et inférieure (UC et LC) sont d'après McLennan [2001]. Les rapports Th/Ta de chaque droite de corrélation sont reportées. (d) Diagramme de corrélation εNd_i en fonction du rapport (La/Nb). La légende des échantillons est la même que pour la figure 6 et les carrés gris clairs supplémentaires de la figure 7c correspondent aux autres tholéiites analysées du Zanskar-Spiti.

(~400 km) suggests a rapid and voluminous effusion of magmatic melts derived from a high degree of partial melting. Such a scenario was already proposed for other Permian tholeiitic series [e.g. Arunachal Pradesh, Bhat, 1984; Bhat and Ahmad, 1990; Central Nepal, Garzanti *et al.*, 1999; fig. 1b], where melt ascent was more efficient through a rifted and strongly thinned lithosphere [e.g. Bhat, 1984; Garzanti *et al.*, 1999].

Compared to the Carboniferous-Early Permian climax of the continental rifting and the associated punctual mainly alkaline magmatism [Pogue *et al.*, 1992; Vannay, 1993; Garzanti *et al.*, 1996a, 1996b; Garzanti *et al.*, 1999], the Panjal Traps extrusion represents a huge magmatic episode characterized by a higher degree of partial melting. Moreover, their rapid effusion could be associated to the drastic lithospheric thinning during continental break-up.

Throughout the Arabian and Indian contiguous continental margins, the available lithostratigraphic informations reported on the southern Neotethyan platforms document that the Neotethyan rifting derived from a long term and probably polyphased tectonic process. In the Himalayan zone, rifting began as early as the Early Carboniferous [Vannay, 1993; Garzanti et al., 1996a; Garzanti and Sciunnach, 1997]. The Lower to Middle Permian Panjal traps extrusion would be one of the largest remnants of the final rifting stage, which accompanied the Neotethyan opening. This huge magmatism belongs to a set of Permian within plate volcanic sequences [fig. 1b], which are systematically regarded as the testimony of an advanced stage of the lithospheric streching [Bhat and Ahmad, 1990; Stampfli et al., 1991; Veevers and Tewari, 1995; Garzanti et al., 1999; this paper]. Syn-rift volcanic successions were also found along the preserved fragment of the neighbouring Arabian continental margin in Oman [Béchennec et al., 1991; Maury et al., 2003; Lapierre et al., 2004]. In Oman, tholeitic and alkaline melts erupted both on the rifted platform (Saih Hatat area, Oman mountains) and in the Hawasina pelagic basin. An OIB-type mantle reservoir has been evidenced, on the basis of Nd and Pb isotopes, beneath the Arabian rifted margin during Middle Permian [Lapierre et al., 2004]. Arabian Middle Permian volcanics yield isotopic compositions, which encompass the signature of the present Panjal basalts

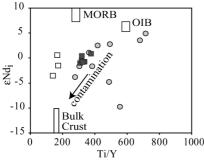


FIG. 8. – ɛNd; vs (Ti/Y) plots of the Panjal Traps from Zanskar-Spiti (Group 1: black squares, Group 2: white squares) and Pakistan (grey circles) [Spencer et al., 1995]. MORB and OIB values are from Sun and McDonough [1989] and Bulk Crust value is from McLennan [2001]. FIG. 8. – Variation des valeurs ɛNd; en fonction de (Ti/Y) des trapps de Panjal du Zanskar-Spiti (Groupe 1: carrés noirs, Groupe 2: carrés blancs) et du Pakistan (ronds gris) [Spencer et al., 1995]. Les valeurs des MORB et OIB sont d'après Sun et McDonough [1989] et celles de la croûte continentale, d'après McLennan [2001].

data. So the Arabian margin can be tentatively compared to a volcanic-type margin considered as related to the melting of a mantle plume, which would have impinged the Arabian lithosphere during or after the oceanic opening [Maury *et al.*, 2003; Lapierre *et al.*, 2004].

Taking into account the discontinuous and diachronic volcanic provinces erupted along the rifted Indian (fig. 1b) and Arabian margins (between Lower Permian to Middle Permian), the model proposing emission of a huge volume of melt in a very short time does not appear satisfactory for the South Neotethyan margin formation. The lack of pre-rift traps is in favour of the active rifting model, as in the North Atlantic magmatic province and the associated Iceland plume [Saunders et al., 1992; Menzies et al., 2002]. Nevertheless, the emission of voluminous within plate volcanic sequence synchronously to Neotethys opening strongly suggests the presence of an abnormally hot asthenospheric reservoir, which should have produced significant volume of tholeitic melts in relation with overlying lithosphere thinning.

Therefore, the more or less wide volcanic outcrops or traps associated to the southern Neotethyan margins rifting could be regarded either as the remnant of independent geodynamic processes suggesting upwelling of several hot spots along the rift [as in Oman, Lapierre *et al.*, 2004; Maury *et al.*, 2003], or as a common geodynamic process related to a passive-type oceanic opening. In this scheme, the triple-point device of the Indian-Arabian-Madagascar Neotethyan rift arms [Stampfli *et al.*, 1991; Veevers and Tewari, 1995; Vai, 2003] could have played a significant role for the localization of melting zones between Indian and Arabian margins.

In the Central Atlantic magmatic province, alternative mechanisms to the mantle plume are actually largely debated. The CAMP represents the wider discontinuous volcanic province related to Pangea fragmentation. It is mainly considered as a passive rifting associated volcanism and not a plume magmatic province [Mc Hone, 2000; Salters *et al.*, 2003; DeMin *et al.*, 2003; Puffer *et al.*, 2003]. An alternative model indicates that wide volumes of basaltic melts can be associated to craton edge-driven convection developed in an abnormally hot upper mantle [Mc Hone, 2000; De Min *et al.*, 2003]. Global warming of the mantle should be itself related to the Pangea supercontinent agglomeration, according to the model proposed by Coltice *et al.* [2007].

Such mechanisms should allow to fit the different Neotethyan continental flood magmatism with a passive rifting model and yield, in particular, an alternative model to the weakly constrained upwelling of a mantle plume set along the same margin. Nevertheless, the geochemical features of Arabian and Indian volcanics discussed in this paper and in Lapierre *et al.* [2004] do not reflect participation of a continental lithospheric mantle component as it is globally accepted for the CAMP related magmas.

Isotopic analyses performed on the entire Panjal Traps and on the Lower Permian tholeiitic sequences along the Indian margin [Garzanti *et al.*, 1999] should allow to better constrain the global geodynamic mechanisms associated to the South Neotethyan margin volcanism. This particularly should highlight a potential participation of the lithospheric mantle, which could represent an alternative OIB-type isotopic reservoir relatively to the mantle plume.

CONCLUSION

This study highlights the geochemical and isotopic characterization of the greatest volcanic episode associated to the formation of the southern Neotethyan margin. This episode led to the eruption of the Panjal Traps on the NW Indian rifted platform. The SE Zanskar-NW Spiti remnants of the Panjal lavas show features of tholeiitic low-Ti (<1.6%) continental flood basalts with LREE, Th enrichments and Nb-Ta negative anomalies (1.0<La/Nb<1.8). The low ɛNdi values (-3.6 to +0.9), and high Pb-isotopic ratios indicate that the Panjal Traps derived from an OIB-like mantle, which was contaminated by an EM II-type component.

- The high La/Nb and Th/Ta ratios combined with the negative ϵNd_i values suggest this enriched component should be the upper continental crust. The crustal contamination process did not occur during an AFC-type process but more probably in the most primitive high-temperature tholeites.
- Trace elements and isotopic ratios also suggest that the Panjal lavas derived from high partial melting degrees of an OIB-like mantle source. The absence of marked Eu anomalies combined with the low proportion of highly fractionated melts supports that Panjal lavas underwent no important differentiation process in shallow magmatic chambers.

– The geochemical features of the Zanskar-Spiti basaltic flows appear largely similar to the coeval Panjal volcanic sequences (Ladakh, Kashmir and Pakistan).

These features allow us to consider the Lower to Middle Permian magmatic event as a melting episode of huge magmatic volumes, generated by a high amount of partial melting in a within-plate setting. Relatively to the Carboniferous to early Permian rifting stages and associated scarce and mainly alkaline magmatism, these Lower to Middle Permian tholeitic melts are regarded as the remnants of the late stages of lithospheric thinning and probably as the record of continental break-up. All these features support geological evidences that the Panjal Traps were extruded during the final step of the Neotethyan rifting episode. In such a scheme, the Panjal Traps were more probably formed in a passive-type rifting environment.

Despite a slight diachronism, similar conclusions are emphasized for the tholeiitic series of the central and eastern segments of the Indian margin, and in the adjacent Arabian margin. Whereas mantle plume upwelling is proposed for the Arabian margin volcanism, such a geodynamic model is not applicable to the entire volcanic sequences of the southern Neotethyan margin.

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