



Article

A Remote-Sensing-Based Alteration Zonation Model of the Duolong Porphyry Copper Ore District, Tibet

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Abstract: Remote sensing (RS) of alteration zones and anomalies can provide information that is useful for geological prospecting and exploration. RS is an effective method for porphyry copper mineral exploration and prospecting prediction. More specifically, the Advanced Spaceborne Thermal Emission and Reflection radiometer (ASTER) data, which include 14 spectral channels from visible light to thermal infrared, are useful in such cases. This study uses visible-shortwave infrared and thermal infrared ASTER data together with surface material spectra from the Duolong porphyry copper ore district to construct an RS-based alteration zonation model of the deposit. In this study, an RS alteration zoning model is established based on ground-spectral alteration zoning results. The methods include PCA (Principal Component Analysis), Ratio, and Slope methods. The information obtained by each method is different. RS-based alteration zonation is developed based on the intersection of maps, resultant from the different methods for extracting information related to different minerals. The alteration zonation information extracted from ASTER RS data is consistent with geological observations. Using information from the RS-based model, we mapped the alteration minerals and zones of the Duolong ore district, thereby identifying prospecting target areas of the deposit.

Keywords: porphyry copper deposit; alteration zone; Tibet; ore prospecting; target area; ASTER



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1. Introduction

Multi-spectral remote sensing (RS) techniques have been increasingly and successfully applied in geological prospecting. Early work in this area includes the recognition of limonite alteration using Landsat multi-spectral data [1], as well as the extraction of the spectral characteristics of hydroxyl minerals in the visible near-infrared (VNIR) to short wave infrared (SWIR) bands [2,3]. Since the early studies, multi-spectral RS and allied techniques have been widely used to investigate the presence and type of alteration [4–6]. For example, a principal component analysis (PCA) has been applied to help classify and establish the distribution of alteration minerals in greenstone belt, Minas Gerais, Brazil [7]. Besides PCA, other techniques for extracting remote-sensing-based alteration information include the ratio method, spectral angle mapper (SAM), minimum noise fraction (MNF), and linear spectral unmixing (LSU) [8–12].

RS multi-spectral applications in mineral exploration have been summarized [13,14] and focus on mapping mineral resources and lithological units [14,15], as well as aspects related to RS-detected anomalies, such as deposit type [16], volcanic genesis [17], porphyry copper and epithermal gold exploration [18], and the alteration characteristics of rocks [19]. Landsat TM/ETM (Thematic Mapping /Enhanced Thematic Mapping) data can extract the information of hydroxyl minerals, carbonate minerals, and iron oxides, but cannot be used to distinguish minerals or mineral assemblages such as minerals consisting of magnesium hydroxyl and aluminium hydroxyl, due to band limitations. However, the Advanced Spaceborne Thermal Emission and Reflection Radiometer (ASTER) remote sensor provides

high-spectral-resolution RS data, containing 14 bands, of which 9 bands are VNIR–SWIR RS and the remaining 5 bands are thermal infrared (TIR) RS data. Since the first use of ASTER, some alteration minerals have been identified using VNIR and SWIR RS data, including kaolinite, dickite, illite, and muscovite data for Cerro La Mina, Argentina [20]. Mars and Rowan [21] studied hydroxyl alteration mineral zonation of a porphyry copper deposit, using ASTER VNIR and SWIR data. ASTER-based mineral mapping has also been applied to investigations of geothermal resources [22], iron ore occurrence [23], and gold deposits [24]. ASTER RS data can also be used to perform alteration mineral mapping [25]. An ASTER spectral database containing single mineral and rock spectral information in the range of 400–15,400 nm has been established [26]. ASTER data have also been used to distinguish single minerals, information for which is held in an ECOSTRESS library [27,28]. ASTER TIR data have been applied to mineral mapping and quartz index research, according to the spectral characteristics of various minerals and the thermal infrared characteristics of silicified alteration zones [29], as well as information extracted from quartzite and carbonate rock [30]. Information on mafic–ultramafic rocks has also been extracted using ASTER TIR [31].

Most research on Copper (Cu) deposits using RS focus on extracting the alteration information of altered minerals and iron oxides. Using the spectral characteristics of altered minerals, various methods for extraction. The alteration information extracted by ASTER is mainly aluminum hydroxyl (Al-OH) and magnesium hydroxyl (Mg-OH) [32–35], and most of the research on porphyry Cu deposits using ASTER focus on hydrothermal alteration mapping [36–41]. Recently, a very large porphyry Cu deposit, the Duolong ore district, was discovered in north Tibet. This deposit is the largest Cu deposit in the Bangonghu–Nujiang metallogenic belt and shows highly developed alteration zonation [42,43]. Here, we use ASTER RS data to extract and analyze alteration zonation information for this porphyry Cu deposit, with the objective of guiding mineral prospecting in the region. Using RS methods to establish alteration zonation of porphyry copper deposits in the Bangonghu–Nujiang metallogenic belt should be useful for establishing mineral prospectivity in this region. We focus on alteration zone mapping based on an ASTER RS spectral analysis of surface materials and rocks to extract alteration zonation information and then construct an RS-based model of mineral alteration zonation.

2. Geological Setting

2.1. Tectonic Setting

The Duolong district comprises two major deposits, namely Duobuza and Bolong. The Duolong district is located on the southern margin of the Qiangtang terrane, to the north of the Bangong–Nujiang suture zone (herein after referred to as BNS) in Tibet (Figure 1a). It extends over 2000 km across the central Tibetan plateau and western Yunnan province, and is thought represent the remnants of the Bangong–Nujiang Tethyan Ocean [44].

The oldest rocks in the Duolong district are represented by the Triassic Riganpeicuo Group, which mainly consists of marble and limestone (Figure 1b). Overlying the Triassic rocks is the Jurassic Quse Formation, comprising two units: the basal Quse I unit of quartz sandstone, overlain by the upper Quse II unit that is composed of sequences of intercalated sandstone and slate. Cretaceous Meiriqie Formation is composed up of basaltic andesite and andesite, and widely distributed in the Duolong district (Figure 1c). An extensive emplacement of volumetrically small intermediate–felsic porphyritic intrusions that comprise similar major minerals (quartz, \pm plagioclase, \pm amphibole, \pm biotite) and accessory minerals (magnetite, \pm zircon, \pm apatite) took place from 120 to 110 Ma [45,46], which was accompanied by large-scale mineralization in the Duobuza and Bolong deposits.

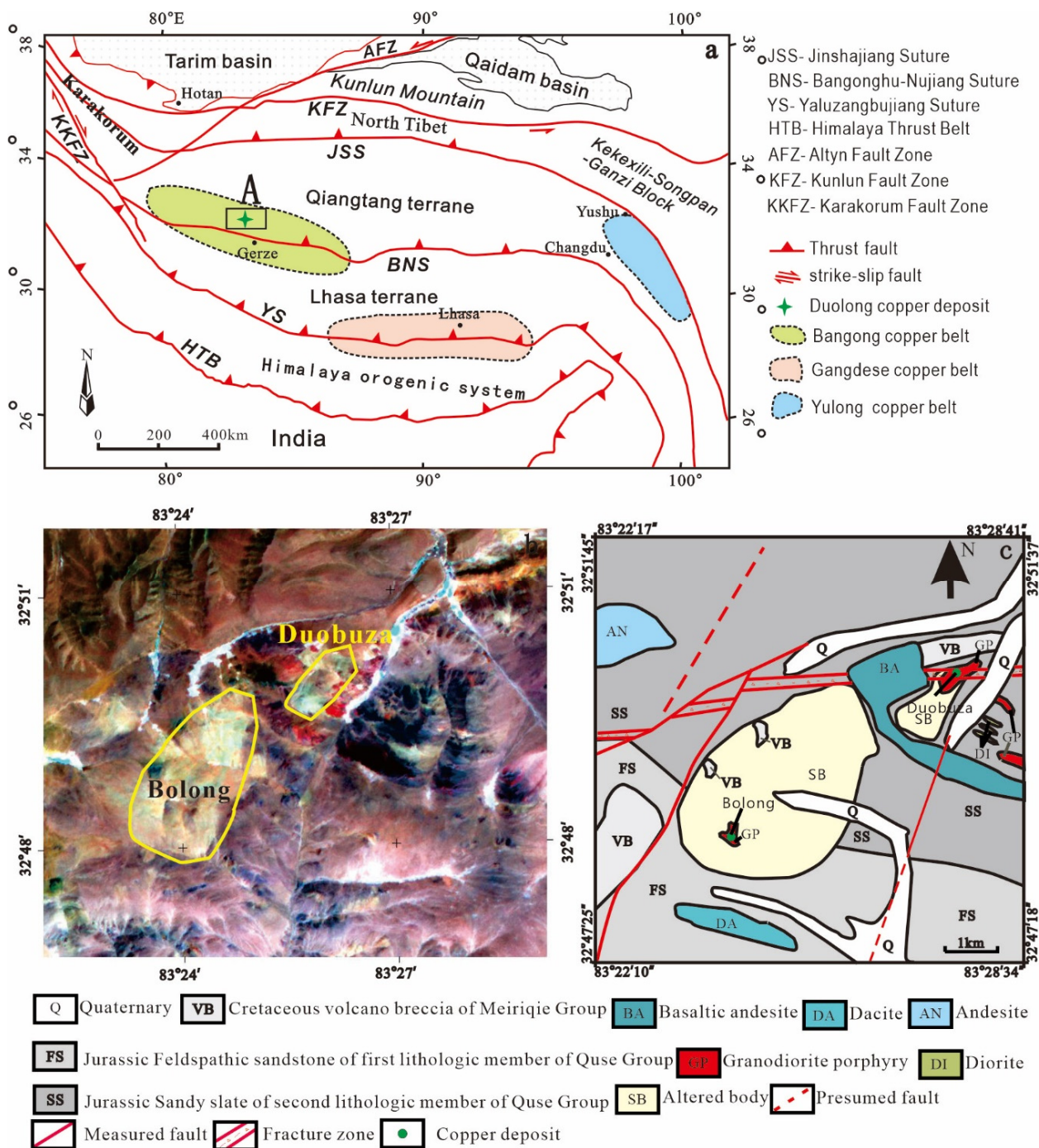


Figure 1. The geotectonic areal division sketch map of Duolong porphyry copper ore field (a); an ASTER false-color composite image of b6(R), b3(G) and b1(B) (b); the geological sketch of Duobuza and Bolong deposits [43] (c).

2.2. Ore Geology

Porphyry Cu deposits are normally located in the basement beneath a comagmatic volcanic pile [47]. Generalized alteration mineralization zoning patterns show that sericitic alteration may project vertically downward as an annulus, separating the potassic and propylitic zones as well as cutting the potassic zone. Sericitic alteration tends to be more abundant in porphyry Cu–Au deposits, whereas chlorite sericite alteration develops preferentially in porphyry Cu–Au deposits. Alteration-mineralization in lithocap is commonly far more complex [48,49]. The Duobuza deposit protrudes in the center of the Duolong district. Several porphyritic intrusions have been recognized based on field mapping and

exploration drilling, two of which are host rocks for Cu–Au mineralization of the Duobuza deposit, a granodiorite porphyry and a quartz diorite porphyry. Hydrothermal alteration and mineralization patterns of the Duobuza deposit are typical of Cu–Au porphyry deposits around the world [50,51]. The mineralized intrusions are affected by an early potassic alteration that is evident from the abundance of magnetite, K-feldspar, biotite and quartz, followed by chlorite alteration and characterized by partial to complete replacement of early potassic alteration minerals by chlorite and minor muscovite. Late sericitic alteration occurs along structurally controlled zones and/or as irregular sericite–quartz–pyrite replacement in altered zones. Most of the ore-grade Cu–Au mineralization at Duobuza appears to have a spatial association with the potassic alteration, which is represented by chalcopyrite, bornite and magnetite-bearing quartz veins as well as disseminations of chalcopyrite, bornite and magnetite.

3. Methodology

3.1. Remote Sensing Data

Landsat TM/ETM data and ASTER data can be analyzed to obtain alteration zone information from porphyry Cu deposits, including the Duolong ore district. TM/ETM data from the Landsat sensor encompass six spectral bands, one panchromatic band, and one thermal infrared band. ASTER data from the TERRA encompass nine spectral bands, one rearview band, and five thermal infrared bands. ASTER data have a wide spectral coverage, high spectral resolution, and are able to provide information for various fields of application. For geological applications, owing to the higher number of shortwave infrared bands (six) and thermal infrared bands (five) of ASTER compared with TM/ETM, ASTER data can be used to distinguish various alteration minerals and mineral assemblages and to examine alteration zonation. To obtain accurate spectral data, we used a field-portable shortwave infrared spectrometer (Analytical Spectral Devices) and a portable infrared mineral analyzer (ASD, Malvern Panalytical Ltd., Malvern, UK).

For this study, spectra of samples obtained from the ground surface were collected using the FieldSpec3 instrument, with a wavelength range of 350–2500 nm. FieldSpec3 has two fast-scanning and reflecting holographic diffraction gratings. Its spectral resolution is 3 nm at 350–700 nm and 10 nm at 700–2500 nm. The analysis of alteration minerals was performed using spectral measurements of surface samples in the horizontal and vertical directions. ASTER data were bought from Pasco China Corporation, Beijing, China in 2008. The scene number is ASTL1B 0704140516580707100089. The data acquisition date is 14 April 2007. The spectral features and variations of alteration zonation were analyzed based on the data collected using the portable instrument. According to these characteristics, ASTER data were used to obtain alteration zonation information for the porphyry Cu using PCA (Principal Component Analysis) and Ratio methods based on the reflectivity of ASTER different wavebands [27,52–56].

3.2. Technical Flow

Alteration minerals and alteration zones are determined according to the spectra of ground samples. ASTER data was selected as the RS data source. The values of each band of ASTER data were compared according to the spectral alteration zonation of ground samples. Based on altered mineral assemblages in different zones, different information extraction methods were designed for each mineral. Ratio, slope, correction absorption, and PCA were used to extract mineral anomalies, and these anomalies were combined to extract alteration zonation. Finally, an RS alteration zoning model was built to interpret multiple hydrothermal centers in Duolong ore district (Figure 2).

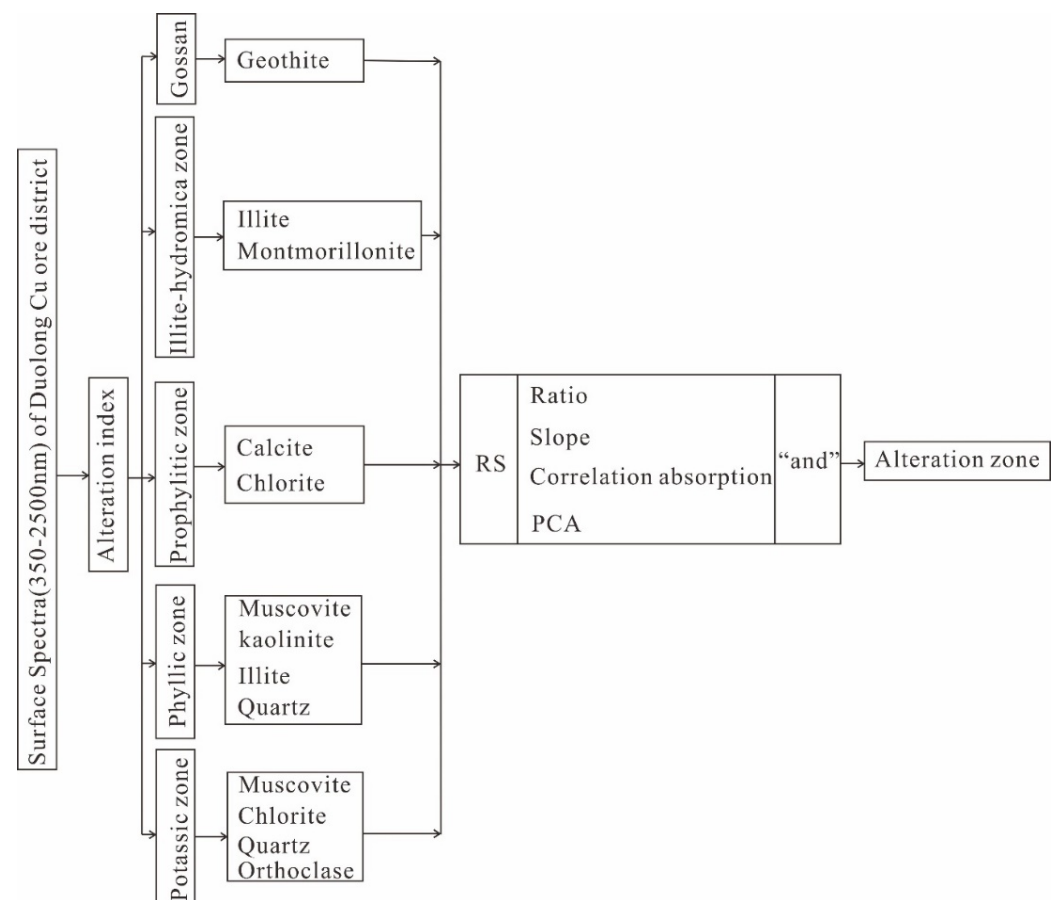


Figure 2. The work flowchart of RS alteration zonation in Duolong porphyry copper ore district.

3.3. Spectral Analysis of Ground Samples

Spectra were collected using a FieldSpec instrument. PimaView [57] software was used to process the spectra. To investigate surface alteration zonation, 280 samples were collected. Each sample was measured three times using the FieldSpec instrument, and each spectrum was then analyzed for alteration minerals using PimaView software.

To obtain surface samples at the Duolong district, we collected metamorphic feldspathic quartz sandstone, granite porphyry, lithic quartz sandstone, gabbro, breccia, granite, basaltic andesite, and quartz porphyry. Of these, 112 samples from the Duobuza deposit and 168 samples from the Bolong deposit were sampled at 300 m intervals according to variations in lithology and alteration (Figure 3).

Spectral reflectances of typical rock samples for each alteration zone, collected from the land surface, were measured using an analytical spectral device (ASD) spectrometer with 2151 bands (350–2500 nm) in the field. The environments of all spectral measurements had sunny and dry weather conditions to reduce the influence of water vapor as far as possible. A contact probe with an internal light source was used to obtain spectra. Dark current correction and white reference correction were made again before each measurement at a new location (Figure 4a). All spectra were also resampled to match the ASTER VNIR–SWIR bands (Figure 4b). Location of the samples is shown in Figure 3. The spectrum number (No.) 2-3 and No. 3-5 were collected in the illite-hydromica zone; No. 8-6 and No. 1-10 were collected in the potassium zone; No. 7-5, No.7-6 and No.7-7 were collected in the propylitic zone; No. 8-8 and No.5-10 were collected in the gossan.

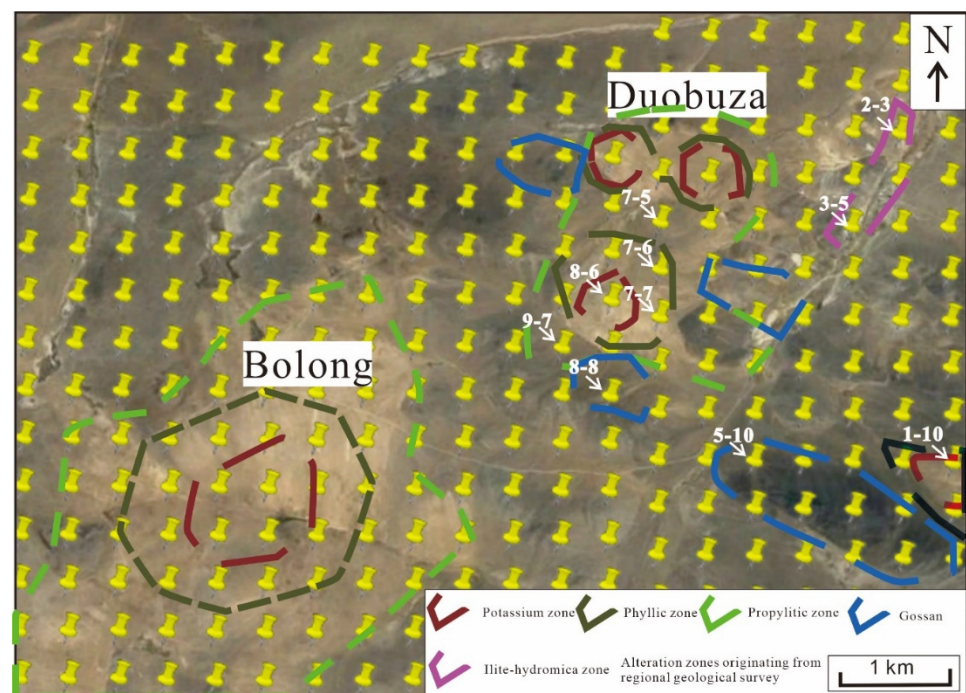


Figure 3. The horizontal alteration zoning of Duolong district.

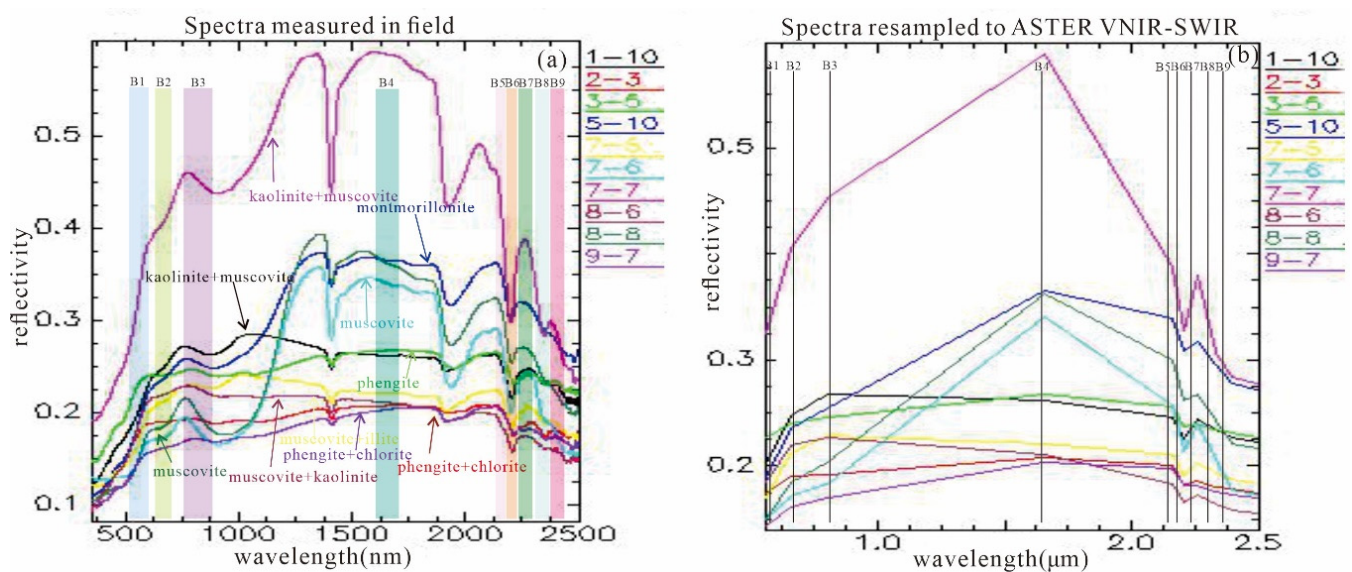


Figure 4. Reflectance curve of typical samples collected from each alteration zone (a) and the spectra curves resampled to the ASTER VNIR–SWIR bands (b).

3.4. Remote Sensing of Alteration Minerals

The major alteration zones of the Duolong ore district include potassic silicification, phyllic, propylitic, illite–hydromica, and gossan zones [17,58]. Alteration minerals vary according to zone, as follows: (i) orthoclase, muscovite, chlorite, and quartz in the potassic silicification zone; (ii) muscovite, kaolinite (for substitute of halloysite), illite, and quartz in the phyllic zone; (iii) calcite and chlorite in the propylitic zone; (iv) illite and montmorillonite in the illite–hydromica zone; and (v) goethite in the gossan zone. The spectral characteristics of all alteration minerals were analyzed over the SWIR and TIR bands and then resampled to ASTER spectral responses [27–53,56] (Figure 5). The RS characteristics of these minerals are provided in Table 1.

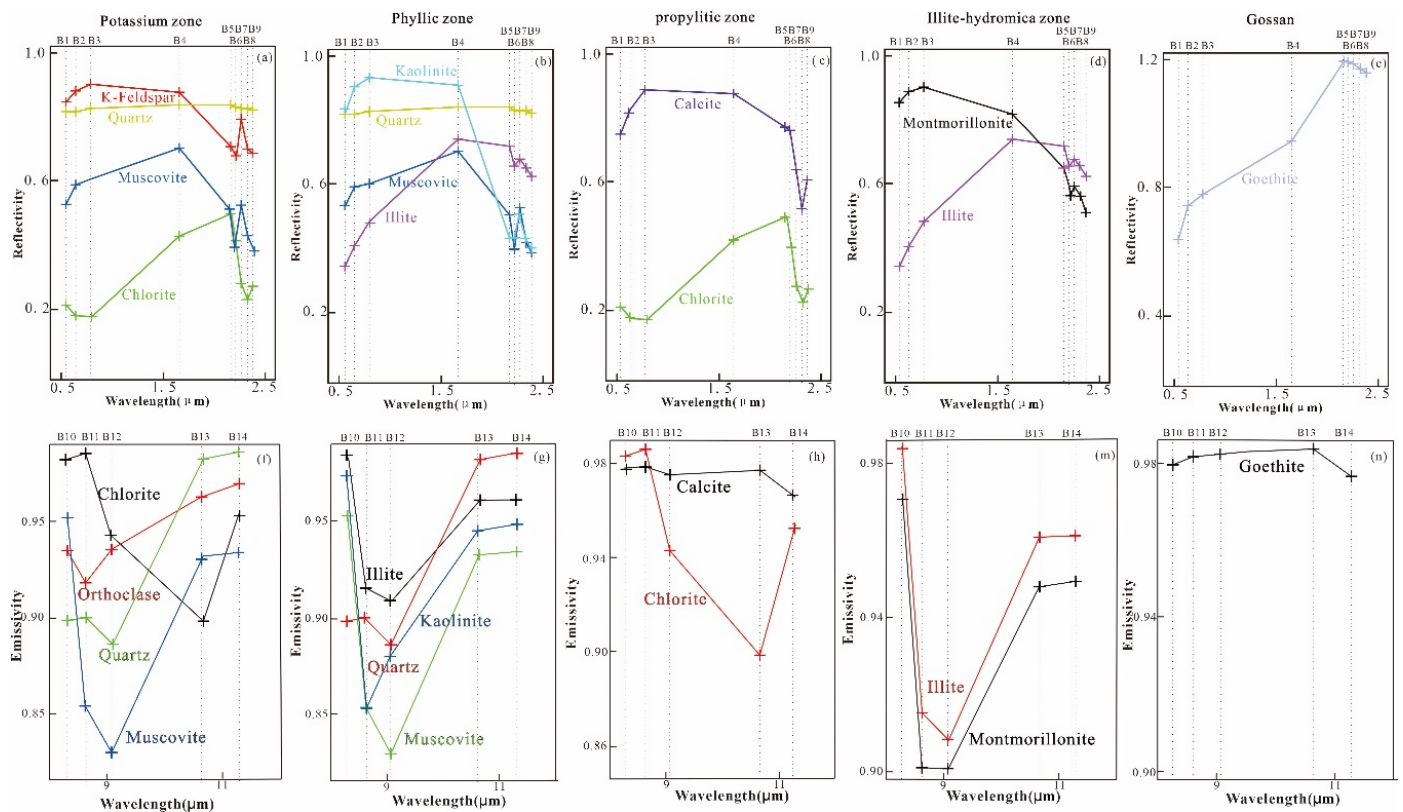


Figure 5. Visible-shortwave infrared and thermal infrared characteristics of each zone of porphyry Cu deposit (resampled to ASTER) [48–52]. Note: The spectra of K-feldspar, Halloysite, and Limonite from the USGS spectral library and wavelength in units of μm .

Table 1. The characteristics of the alteration minerals.

Zones	Bands	ASTER Bands													
		VNIR				SWIR					TIR				
		B1	B2	B3	B4	B5	B6	B7	B8	B9	B10	B11	B12	B13	B14
Potassium silicification zone	Orthoclase	H	H	H	L	L	H	L	L	-	L(L)	L	H	H	
	Muscovite	L			H	L	H	L	L	-	L	L	H	H	
	Chlorite	L			H	H	L	L		H	H	-	L	-	
	Quartz	H			H	H				L	L	L(H)	L	H	
Phyllic zone	Mucovite	L			H	L	H	L	L	-	L	L	H	H	
	kaolinite				H		L	L	L	-	L	L	H	H	
	Illite				H	L			L	-	L	L	H	H	
Propylitic zone	Quartz	H			H	H			L	L	L(H)	L	H	H	
	Calcite				H	H			L	H	H	H	H	L	
Illite-hydromica zone	Chlorite	L			H	H			L	L	H	H	-	L	
	Illite				H	L			L	-	L	L	H	H	
Gossan	Montmorillonite				H	H	L	L		L	L	-	L	L	
	Goethite	L			H					L	H	H	H	L	

Note: “H” indicates a high value of reflectance or emissivity. “L” means a low value of reflectance or emissivity. VNIR: The Visible and Near-Infrared; SWIR: The Short-Wave Infrared; TIR: The Thermal Infrared.

3.5. Alteration Zones Information Extraction

According to the characteristics of alteration minerals (Table 1), we used a variety of techniques to analyze spectral information, including PCA, the ratio method, the slope method, correlation absorption, polynomial equations, and the thermal and visible-shortwave infrared data of ASTER. Among these methods, PCA and ratio are quoted from Ninomiya and Crosta et al. [7,54,55]. Additionally, slope and correlation absorption are from the ratio, in order to amplify the spectral characteristics of minerals. For the same kinds of minerals, relevant information can be extracted by one method, which covers a large range. By designing these methods to extract the intersection information, the results are closer to the actual situation. Table 2 summarizes the main methods of information extraction with respect to the spectral characteristics of each alteration mineral. Different thresholds are used for each method in order to obtain a more suitable alteration range. The results for each method are shown in Figure 6.

Table 2. The algorithm of information extraction of each alteration mineral of the Duolong ore district.

Minerals	Methods	Ratio	Slope	Correlation Absorption	PCA [7]
Quartz		$B11 \cdot B11 / B10 / B12$ [55]			
Chlorite		$B5 / B8$	$(B5 - B8) / B5$	$(B4 - B9) / (B4 - B8)$	PCA1.3.4.8 PC4
Orthoclase		$B7 / B6$ $B11 \cdot B11 / B10 / B12$ $B12 \cdot B14 \cdot B14 \cdot B14 / B13 /$ $B13 / B13 / BA13$ [54,55]	$(B4 - B6) / B4$	$(B4 - B6) / (B4 - B7)$	PCA1.3.4.6 PC4
Muscovite		$B4 / B6$	$(B4 - B6) / B4$	$(B4 - B6) / (B4 - B7)$	PCA1.3.4.6 PC4
Kaolinite		$B4 / B6$	$(B4 - B6) / B4$	$(B4 - B6) / (B4 - B7)$	PCA1.3.4.6 PC4
Calcite		$(B6 + B7) / (2 \cdot B8)$ $B13 / B14$ [54,55]	$(B4 - B8) / B4$	$(B4 - B8) / (B4 - B6)$	PCA1.3.4.8 PC4
Illite		$(B7 + B5) / (B6 + B9)$	$(B4 - B6) / B4$	$(B4 - B9) / (B4 - B6)$	PCA1.3.4.9 PC4
Geothite		$B3 / B6$			
Montmorillonite		$(B7 + B5) / (B6 + B9)$	$(B4 - B6) / B4$	$(B4 - B6) / (B4 - B7)$	PCA1.3.4.6 PC4

Note: B1 is first band of ASTER data. PCA: Principal Component Analysis. Some algorithms are given in experimental formula.

3.6. Application of Magnetic Anomalies

Hydrothermal alteration causes mineralogical changes in the host rocks of ore bodies, and consequently, the original magnetic properties of these rocks could be strongly modified [58]. In Duolong district, a high-precision magnetic survey was performed. From the magnetic survey results, the porphyry Cu deposits were found to be in low positive magnetic anomalies, of around 300–650 nT [43].

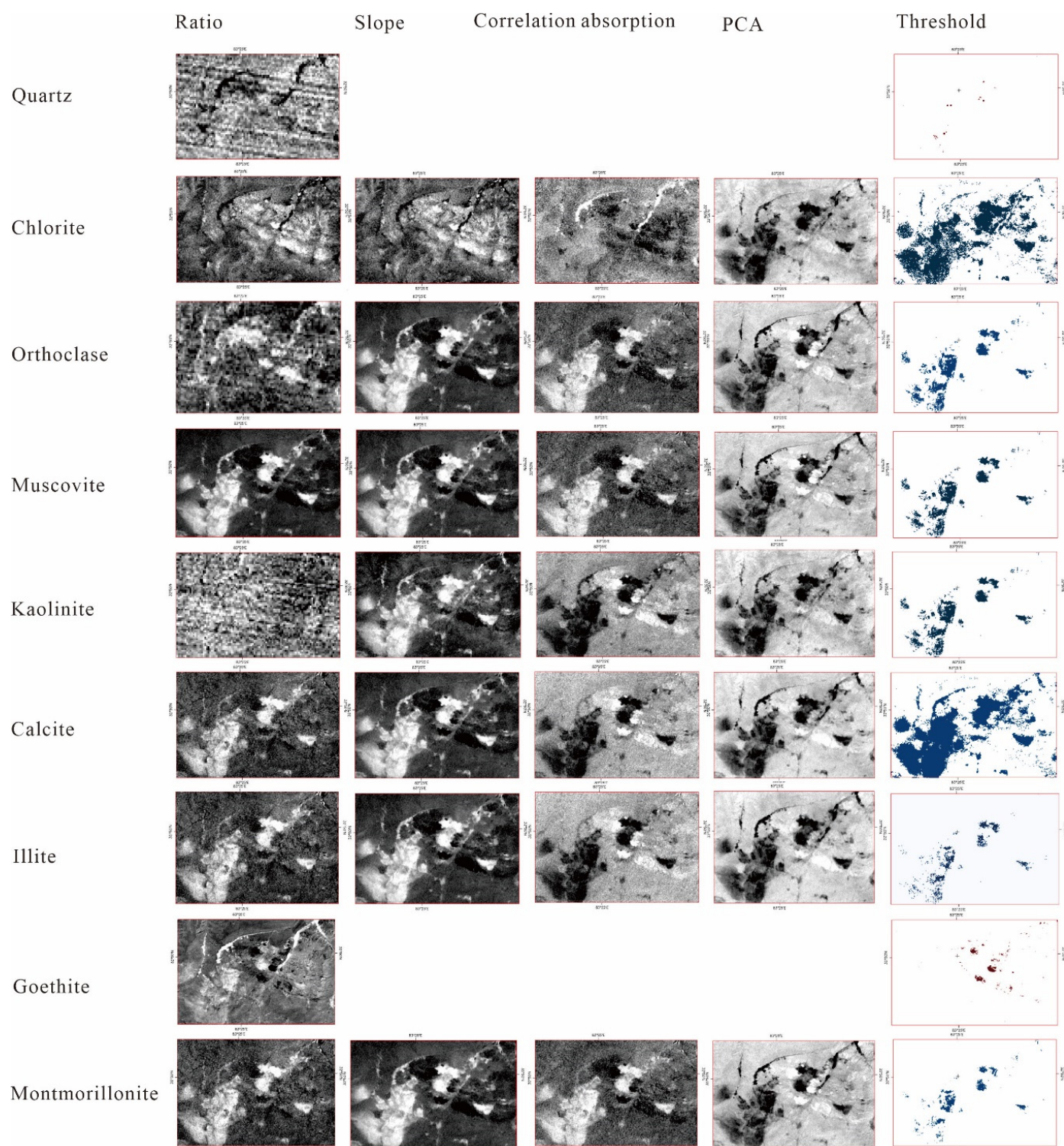


Figure 6. The results of each method and merging information based on threshold.

4. Results and Analysis

According to analysis results of the spectra of surface samples using PimaView software, the alteration mineral assemblage of the Duobuza and Bolong deposits comprises kaolin, white-mica, smectite, chlorite, epidote, carbonate, but 10% of the samples are invalid. The alteration types of surface samples include limonitization, propylitization, chloritization, and sericitization (Figure 7).

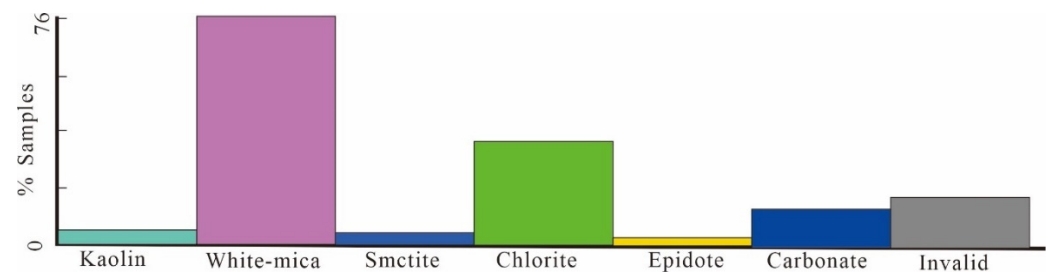


Figure 7. Frequency of alteration minerals interpreted by surface spectra.

4.1. Horizontal Alteration Zonation

According to the frequency of minerals analyzed by spectra (Figure 7), muscovite, chlorite, and calcite were used to divide the alteration zonation with respect to the horizontal dimension because of high frequencies. The distribution of the main alteration minerals in the alteration zones at the Duolong district is as follows: potassic silicification zone: K-feldspar, kaolinite and muscovite; phyllic zone: kaolinite and muscovite; propylitic zone: chlorite and calcite.

The outcome is therefore a semi-quantitative description of the alteration zonation. The obtained results show that the potassic silicification zone is located mainly in the northeast of the Duobuza deposit, whereas the phyllic zone is in the center of the deposit and the propylitic zone is located in the periphery. In the Bolong deposit, the potassic silicification zone occurs in the western part, and the phyllic and propylitic zones are found in the southeastern part (Figure 3).

4.2. Remote Sensing Model

Alteration zonation information was extracted using superposition of minerals corresponding to each zone, including union information extracted using all methods, and revealed the following: quartz, orthoclase, muscovite, and chlorite in the potassic silicification zone; quartz, muscovite, halloysite, and illite in the phyllic zone; calcite and chlorite in the propylitic zone; illite and montmorillonite in the illite–hydromica zone; and limonite in the gossan zone. Results of the RS-based alteration zonation are shown in Figure 8.

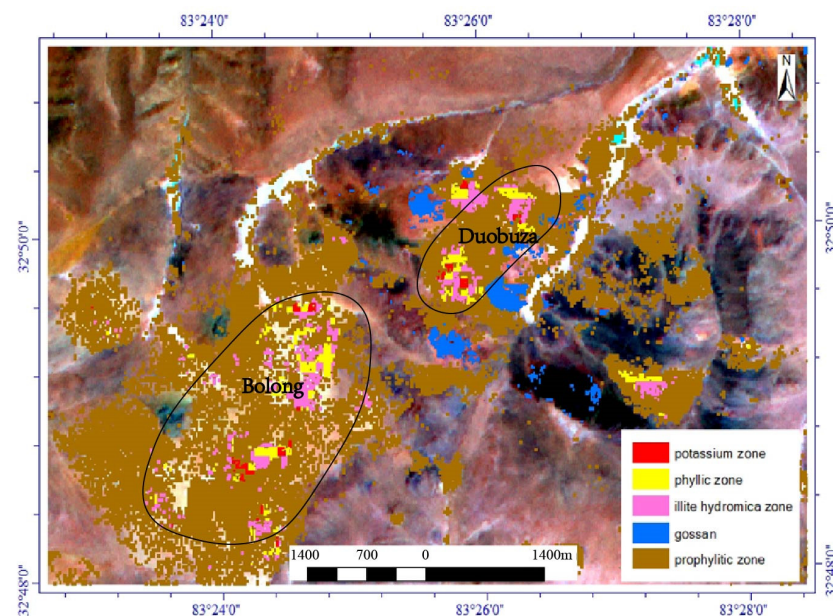


Figure 8. The remote sensing alteration zoning of Duolong deposits extracted from ASTER.

4.3. Field Validation

The alteration zones of Duolong ore district in Tibet were verified in the field (Table 3). The corresponding results of the potassic zone, phyllic zone and propylitic zone as extracted by RS are consistent with field (Figure 9). The main minerals in the potassic zone are quartz and orthoclase, and some kaolinite. The main altered minerals in phyllic zone are muscovite, halloysite, kaolinite and illite. Chlorite and calcite are the main altered components of propylitic zone. These minerals are generally consistent with the altered minerals analyzed by spectra. Additionally, the minerals composition and distribution are basically consistent with the minerals extracted by RS.

Table 3. Field verification points.

Field No.	Longitude	Latitude	Altered Rocks	Alteration
D01	83.45036111	32.84075	Propylitic quartz sandstone	Chloritization, Carbonation
D02	83.44861111	32.83725	Potassic quartz porphyry	Potassium, Silicification
D03	83.44538889	32.83722222	Potassic quartz porphyry	Potassium, Silicification
D04	83.34377778	32.73613889	Phyllic quartz sandstone	Muscovitization, Kaolinization
D05	83.34375	32.73641667	Phyllic quartz sandstone	Muscovitization, Kaolinization
D06	83.34116667	32.74077778	Phyllic quartz sandstone	Muscovitization, Kaolinization
D07	83.33833333	32.74188889	Phyllic quartz sandstone	Muscovitization, Kaolinization
D08	83.33583333	32.74344444	Phyllic quartz porphyry	Muscovitization, Kaolinization
D09	83.33947222	32.73866667	Phyllic quartz porphyry	Muscovitization, Kaolinization
D10	83.29894444	32.72783333	Phyllic quartz sandstone	Muscovitization, Kaolinization
D11	83.39752778	32.807	Propylitic quartz sandstone	Chloritization, Carbonation
D12	83.39363889	32.812	Propylitic quartz sandstone	Chloritization, Carbonation
D13	83.39238889	32.81647222	Propylitic quartz sandstone	Chloritization, Carbonation
D14	83.38902778	32.82333333	Propylitic quartz sandstone	Chloritization, Carbonation
D15	83.38833333	32.81644444	Propylitic quartz sandstone	Chloritization, Carbonation
D16	83.40566667	32.80652778	Potassic quartz sandstone	Potassium, Silicification
D17	83.41097222	32.81058333	Propylitic quartz sandstone	Chloritization, Carbonation
D18	83.44463889	32.83075	Propylitic quartz sandstone	Chloritization, Carbonation
D19	83.43777778	32.8275	Propylitic quartz sandstone	Chloritization, Carbonation
D20	83.44347222	32.83641667	Propylitic quartz porphyry	Chloritization, Carbonation
D21	83.70294444	32.79388889	Phyllic schist	Muscovitization, Kaolinization
D22	83.54788889	32.78622222	Propylitic quartz sandstone	Chloritization, Carbonation
D23	83.56686111	32.78875	Phyllic quartz porphyry	Muscovitization, Kaolinization
D24	83.58688889	32.82641667	Potassic quartz porphyry	Potassium, Silicification

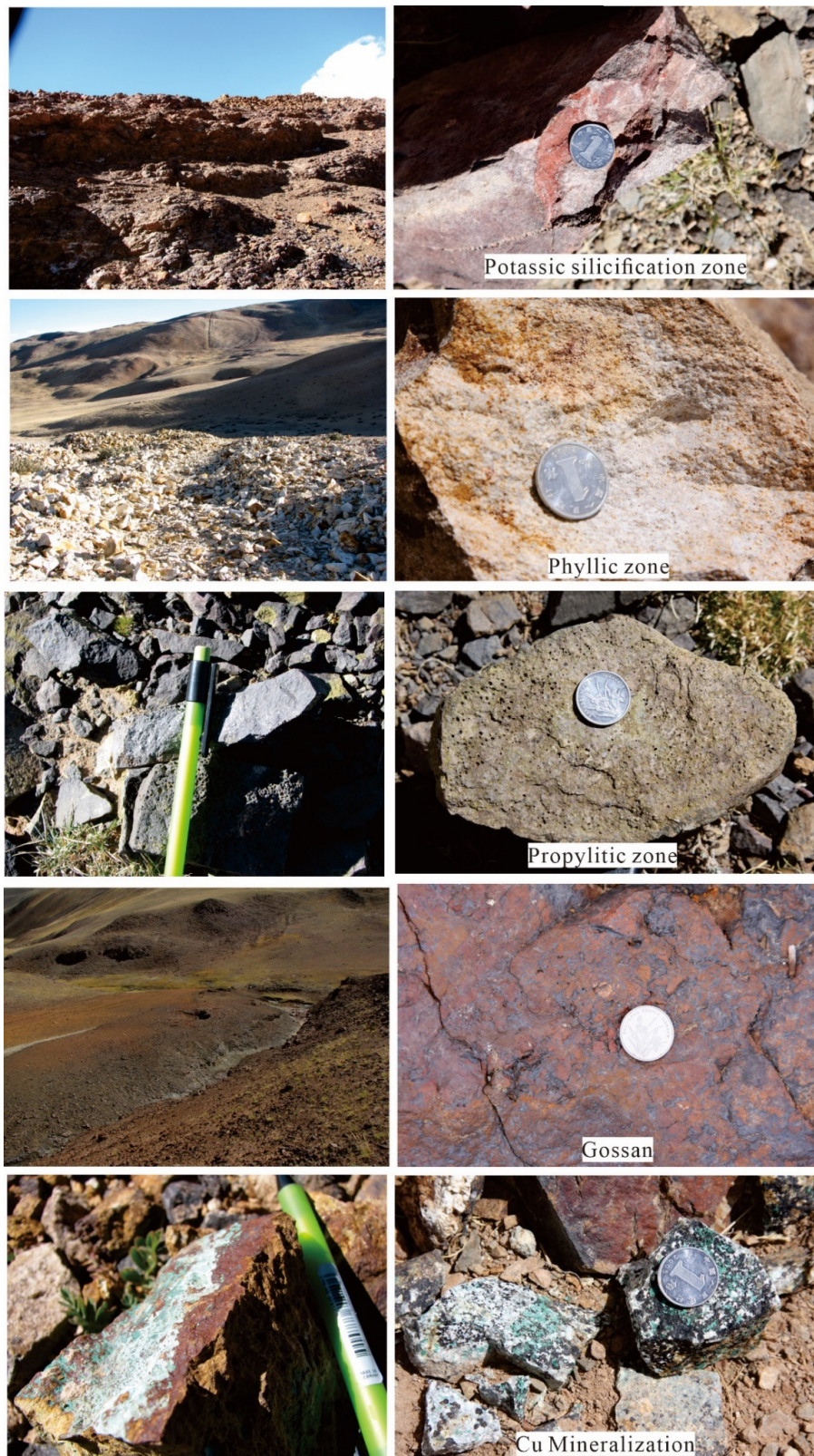


Figure 9. Field validation of alteration zones.

5. Discussion

Porphyry Cu deposits are a veinlet type of sulfide copper deposit formed by a post magmatic medium high temperature hydrothermal solution of potash and hydrogen altered mineral [17,59]. Ideal porphyry Cu deposits are typically related to hydrothermal alteration zones, such as potassic, phyllic and propylitic zones [60]. Hydrothermal alteration minerals with diagnostic spectral absorption properties in VNIR, SWIR, and TIR can be identified by ASTER RS data [17,21,23,26,37,61–66]. The core of the potassium zone is surrounded by phyllic and propylitic zones. The phyllic zone is characterized by sericite, which has an Al-OH absorption feature at 2.20 μm , coinciding with ASTER band 6. The mineral assemblages of the outer propylitic zone include chlorite, epidote, and calcite have absorption features at 2.35 μm , coinciding with ASTER band 8 [2,17,21,63,67–69]. ASTER provides useful data for studying mapping hydrothermal alteration mineral zones associated with porphyry copper. Several spectral mineralogic indices, using ASTER VNIR, SWIR, and TIR bands, were proposed [70–80]. These indices provided accurate spectral information for minerals [17]. The PCA method can extract the characteristic spectral information of key hydrothermal alteration mineral end-members [76]. These methods can be used to extract information of minerals or mineral assemblages.

Previous studies indicated small intermediate–felsic porphyritic intrusions and Cu-Au sulfide ores in Duolong ore district were from 120 to 110 Ma [45,46], which was accompanied by large-scale mineralization. In fact, aluminum hydroxyl (Al-OH) and magnesium hydroxyl (Mg-OH) were important prospecting indicators [36–39]. The advantages of the RS alteration model of porphyry copper deposits mainly correspond with alteration zones of porphyry copper deposits. The study indicated multiple hydrothermal ore-forming centers with obvious signs. The magma involved in the formation of the porphyry copper deposit is inferred to have been intermediate–felsic in composition [38], forming hypabyssal or ultra-hypabyssal rocks and producing altered minerals. The diagnostic spectral absorption features of hydroxyl-bearing minerals including clay and sulfate groups, as well as carbonate minerals due to vibrational processes of fundamental absorptions of Al-OH, Mg-OH, and CO_3 groups in the SWIR and TIR regions of the electromagnetic spectrum [2,17,67,68,77]. Some spectra of surface samples were collected. These spectra indicate spectral features of kaolin, white-mica, chlorite, epidote, carbonate and smectite by PimaView software analysis. These mineral assemblages belong to the alteration minerals of porphyry Cu deposits [60]. According to the combinations of alteration minerals in each alteration zone, the information extracted by RS corresponds to the characteristics of quartz, potash feldspar, kaolin, white-mica, chlorite, epidote, carbonate. In these samples, quartz, potash feldspar, biotite, anhydrite, halloysite and illite cannot be identified. Therefore, when the information of the alteration zones was extracted using RS image, we had to recommend the reference spectra [48–52]. Our ASTER RS-based alteration zonation model for the Duolong ore district should help to guide mineral exploration in the region. The alteration zonation model shows that the magmatic rock mass occupies the center of the deposit, the near periphery comprises the sericite zone, and the distal periphery contains the propylitic zone, with the illite–hydromica zone intersecting across these zones. The gossan zone is located in the periphery of the propylitic zone. During the ascent of the magma that was emplaced in the hypabyssal subsurface in the deposit area, the crust became domed, and fractures developed, providing channels for magmatic–hydrothermal fluids to migrate. The country rock was altered to form the potassic silicification zone. Biotite in the alteration zone was altered to chlorite, and K-feldspar was altered to muscovite and sericite. At a distance from the magmatic body, sericitization and silicification occurred in the surrounding rock outside the potassic silicified zone, with the inner part of this zone being characterized by sericite, illite, kaolinite, and halloysite. Further out, the illite–hydromica zone and the gossan zone constitute the propylitic zone, containing predominantly chlorite and calcite, and the sericitization zone. These zones were largely formed by the replacement and sulfurization of iron-bearing materials in volcanic rocks of the Early Cretaceous Meiriche Formation, by sulfur-containing magmatic–hydrothermal

fluids (Figure 10). Meanwhile, hydrothermal magnetite coexists with chalcopyrite at Duolong ore district, consistent with the mineralization characteristics of porphyry Cu–Au deposit [78–80]. Positive and low magnetic anomalies indicate the location of metallogenic porphyry (Figure 11). Here, we used RS to extract alteration zonation information for the Duolong porphyry copper ore district, based on geological information and the spectra of altered rocks of the deposit, after which we constructed a model of the zonation. Our model can be applied to optimizing ore prospecting target areas in the deposit. The metals and ore-forming fluids of Porphyry deposits in Duolong ore district are more likely to be released from magma chamber [45,46].

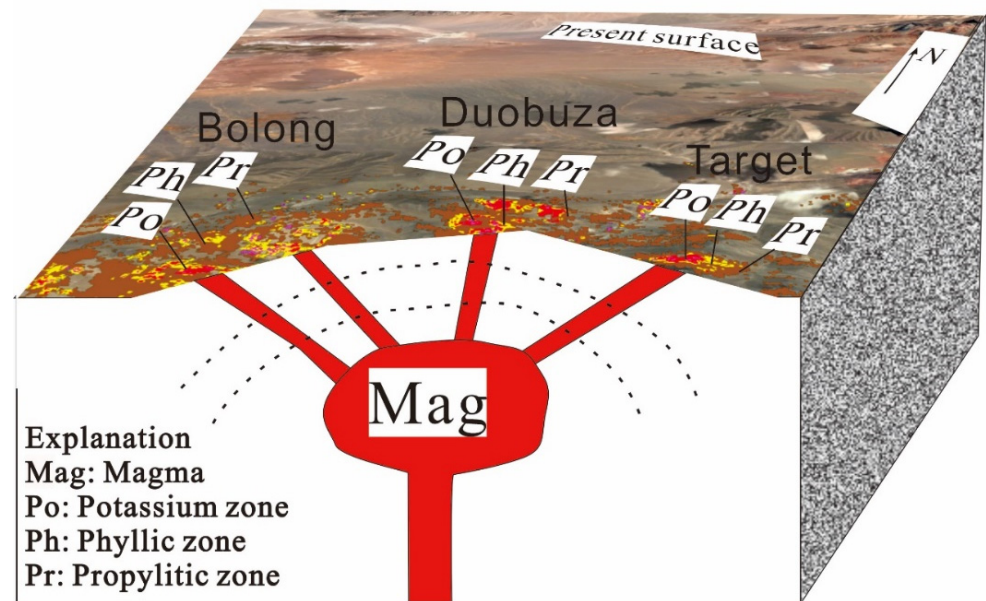


Figure 10. Remote sensing alteration zoning model of porphyry copper deposits.

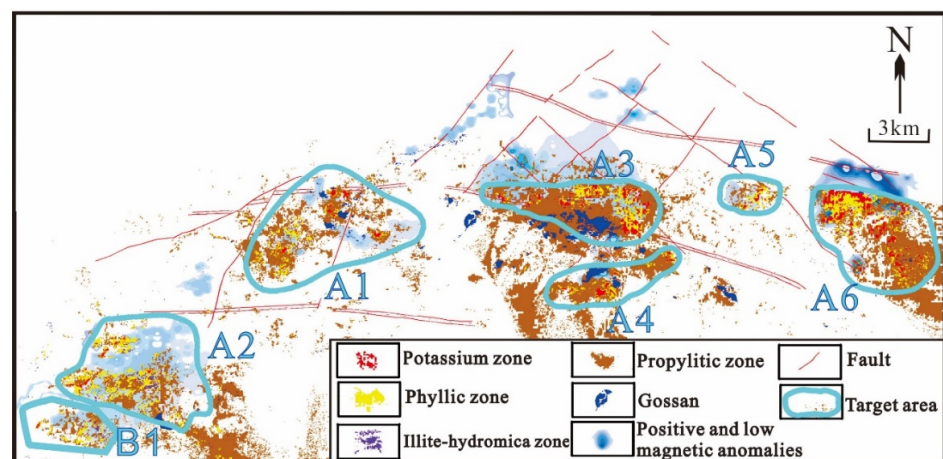


Figure 11. Target area optimization map of Duolong district [43].

The RS-based model of alteration zonation constructed for the Duolong ore district was integrated with various prospecting factors in the western part of the Bangong–Nujiang metallogenic belt. Factors related to prospecting include magmatic rock type, country rock, geological structure, alteration zonation, geophysical properties, and geochemical characteristics [43]. These factors were superposed to evaluate prospectivity using GIS software. The predicted areas were located in typical RS alteration zones, in a favorable metallogenic geological position, and have a high geochemical value area of Cu or Au, as

well as positive and low magnetic anomalies. Seven target areas were selected for porphyry Cu ore-deposit prospecting, distributed along the large-scale structural belt of the Tiegelong Mountains (Figure 11). Of the seven areas, six were class A target areas and one was a class B target area. A class A target area generally has outcrops of granodiorite porphyry and favorable geochemistry, alteration zonation, and magnetic anomalies, whereas a class B target area lacks one or two of these favorable characteristics.

6. Conclusions

We investigated alteration zonation and prospectivity in the Duolong porphyry copper ore district by analyzing the remote sensing (RS) spectra of altered rock on the surface and integrating this information with the deposit geology of the Duolong ore district. We established the spectral characteristics of altered minerals and distinguished alteration zones in the deposit. The mineralization of the deposit is magmatic–hydrothermal in origin and alteration zonation extends from the center of the deposit, where the magmatic body was emplaced into country rock, to the periphery. From the center to the periphery, with decreasing temperatures of magmatic–hydrothermal fluids from medium–high to medium–low, the order of alteration zoning is as follows: (i) potassic silicified zone (containing biotite and chlorite); (ii) sericitization zone (Sericite, illite and kaolinite); (iii) argillization zone (kaolinite, halloysite, chlorite, illite, and montmorillonite); and (iv) propylitic zone (carbonate, chlorite epidote, and montmorillonite).

In this study, an ASTER RS alteration zoning model was established based on ground spectral alteration zoning results. The methods include PCA (Principal Component Analysis), Ratio, and Slope methods. RS-based alteration zonation was developed based on the intersection of maps, resultant from the ratio, slope, extraction information of different minerals obtained through PCA methods. An RS-based alteration zonation model of the Duolong copper porphyry district was constructed by combining alteration-mineral and zone-spectral information with the mineral and geological characteristics of the deposit. This suggests that the model provides a suitable tool for establishing alteration zonation and, when combined with other information, for predicting prospectivity. Combining the RS-based alteration-zonation information with prospecting factors such as magnetic anomaly and geological structure information, allowed target areas to be established for the Duolong ore district. This method is applicable to an exploration of other similar porphyry copper deposits. In future, we will identify minerals using hyperspectral embedded with an unmanned aerial vehicle (UAV). A three-dimensional prospecting method will be built by satellite RS, UAV hyperspectral and ground and boreholes.

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