Multivariate statistical analysis and Geospatial approach for evaluating Hydro geochemical characteristics of meltwater from Shaune Garang Glacier, Himachal
 Pradesh, India

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10 Abstract

11 The study focuses on the hydro-geochemistry of Shaune Garang glacier's meltwater concerning glacial geomorphology. Seventy-nine water samples (53 in 2016 and 26 in 2017) of ablation 12 season were analyzed. The cations were dominant in the order $Ca^{2+} > Mg^{2+} > Na^+ > K^+$, and 13 the anions in the order $HCO_3^- > SO_4^{2-} > Cl^- > NO_3^-$. The result demonstrated that HCO_3^- were 14 the abundant ions, accounting for 41.03 and 34.84 % of the total ionic budget (TZ). The high 15 ionic proportions of $(Ca^{2+}+Mg^{2+})$ versus TZ^+ and $(Ca^{2+}+Mg^{2+})$ versus (Na^++K^+) were identified 16 17 as the primary factors influencing dissolved ion chemistry in meltwater. Piper diagram shows that Ca^{2+} HCO₃⁻ type water is the most common, followed by Mg²⁺ HCO₃⁻. In addition, a 18 remote sensing approach has been used to find the possible source of the chemical constituents 19 in the meltwater. The catchment geology has been mapped on various scales, including diverse 20 rocks and unconsolidated surface materials containing "quartz and carbonate minerals". 21 Layered silicates (LS) and "hydroxyl-bearing minerals" are not as common as they used to be, 22 but their availability varies greatly in the area where they are found. The distribution of Layered 23 silicate (LS) minerals within the catchment are majorly found at lower altitudes, which implies 24

the weathering mechanism due to the interaction of meltwater and parental rock. Multivariate analysis revealed that CO₃ and SiO₂ weathering, sulphate dissolution, and pyrite oxidation dominate dissolved ion concentrations. Chemometric analysis of meltwater hydrogeochemistry through Principal Component Analysis (PCA) explains 72.1 % of the total variance of four PCs. PCs 1, 2, 3 and 4 explain 39.21%, 12.91%, 10.24% and 9.74% of variance respectively in 2016. Similarly, in 2017, four PCs explain 69.9.1% of the total variance. PC 1, 2, 3 and 4 can explain 26.62 %, 20.12 %, 12.64 % and 10.52 % of variance.

32 Keywords: Hydro-geochemistry, Geological mapping, Chemometric analysis, Shaune Garang
33 glacier, Hydroxyl-bearing minerals.

34 Introduction

The Hindu Kush Himalayan Mountain ranges and the Tibetan Plateau extend across several 35 36 Asian countries (Bolch et al., 2012) and are recognized as the cryospheric region beyond the polar latitudes (Bolch et al., 2019; Wood et al., 2020). The Hindu Kush Himalayas, the source 37 of many essential river systems in Asia, are the highest mountain range (Kaser et al., 2010; 38 39 Singh et al., 2020; Immerzeel et al., 2020), which provide a source of livelihood for millions of people living in this region. The Hindu Kush Himalaya, also known as the 'Third Pole' and 40 the 'water tower of Asia,' provides ecosystem services, particularly water, the lifeblood of all 41 organisms. The ecosystem services produced in the region provide support for 240 million 42 43 people who live in the hills and mountains and approximately 1.7 billion people who live 44 downstream in the major river basins. The increasing metropolitan infrastructure in the Himalayan region in the last three decades has put additional strain on the region's limited 45 water resources (Tiwari et al., 2018). Other than natural forces, the primary causal variables in 46 47 the Himalayas include extraordinary urbanization, increasing tourism, shifting land use patterns, garbage disposal, and agricultural runoff are all factors to consider (Thakur et al., 48

2019; Kumar et al., 2019). Snow/glaciers meltwater is a vital water source for domestic use in 49 the higher Himalayas. At the same time, it is essential for irrigation, industrial use, and 50 51 hydroelectric power generation for the downstream population (Singh et al., 2008). When the other water sources are low in supply, the glacier discharge plays a more prominent role in 52 sustaining the demand (Kumar et al., 2016; Singh et al., 2022). The meltwater drains from the 53 glaciers meet the various soil particles and are subsequently influenced by the various mineral 54 55 components. According to recent research, climate and land-use/cover changes caused by human endeavours significantly impact the chemical composition of Himalayan freshwater 56 57 (Pant et al., 2021; Thapa et al., 2020). Hydrogeochemical characterization of the glacier meltwater varies between glaciers due to differential lithology (Collins, 1979). When it comes 58 to glacier meltwater chemistry and, consequently, the quality of meltwater, the geology of the 59 catchment is one of the most important factors to consider. Such studies also improve our 60 understanding of the interactions of meltwater with the underlying geological strata and 61 provide insights into the hydro-climatic regime of a region. There is a need to conduct more 62 research in these areas to create a robust database to determine hydrochemistry confidently. 63 The widely varying geology makes the hydro-chemical study of Himalayan glaciers even more 64 exciting. Earlier studies on the chemistry of the major ions in the Himalayas engrossed in 65 recognizing chemical input in glacial meltwater. Various studies of Himalayan glaciers have 66 suggested that water transients over sub-glacial waterways encounter rock substrate 67 68 experiencing a significant chemical change in meltwater (Singh et al., 2015, 2017). Some studies specify that the high degree of differential erosion in Himalayan glaciated areas is the 69 consequence of the long interaction time of meltwater through bedrock (Haritashya et al., 2010; 70 Singh et al., 2015). Investigating hydrochemistry in the Himalayas is highly important, and 71 several researchers (Haritashya et al., 2010; Singh et al., 2015; Kumar et al., 2019) have 72 contributed to this regard. 73

All the previous studies in the Himalayan region on water chemistry have focused only on the 74 chemical characterization of meltwater from glaciers. Documentation of such investigations 75 substantially allows precise lithological mapping and erosion rate estimation in these high-76 77 mountain areas. This helps in understanding large-scale links between glacio-hydrology and geochemistry. Considering possible sources and their differential erosion rates can give us a 78 better idea about the possible future trajectory of meltwater chemistry in these high-mountain 79 80 catchments. Photogrammetry, high-resolution terrain modelling and hyperspectral imaging are used in lithological mapping and estimation of erosion rates in high mountains. Further, it helps 81 82 establish large-scale links between glacio-hydrology and geochemistry on a global scale.

In the present study, the remote sensing approach has been used to identify chemical species' 83 sources of origin in meltwater through the catchment scale lithological mapping, mineral 84 detection, and many other perspectives. Also, ASTER-SWIR data were used to delineate types 85 of rocks in the Shaune Garang catchment to interpret and measure minerals and the 86 geochemical composition of debris on the glacial surfaces. This study validates several features 87 of ASTER (SWIR and TIR) data and their processing techniques; analysis has been used to 88 map several minerals containing indices. These exploration approaches are suitable for low-89 cost identification and mapping of minerals containing indices such as layered silicates (LS), 90 Calcite (CA), Hydroxyl bearing (OH), Alunite (AL) on the SWIR band and carbonate, quartz, 91 92 and mafic index on TIR band in ASTER data. In this research, we focus on identifying (i) major 93 ion concentrations, chemical characteristics, and their variability; (ii) hydro-geochemical processes and solute sources during the study period; and (iii) lithological mapping of the 94 Shaune Garang catchment. Hence, the results of this study enrich the catchment findings in 95 terms of glacio-hydrology and geochemistry. Furthermore, this would be a significant step 96 towards a more precise interpretation of weathering and hydrological processes in the 97 catchment. This research includes applying various chemometric analyses such as principal 98

99 component analysis (PCA) and factor analysis to understand better the dominant weathering100 process in the Shaune Garang catchment.

101 Study Area

The hydro-geochemical analysis has been performed in the Shaune Garang glacier catchment, 102 located in Himachal Pradesh. The glacial meltwater of this catchment merges to the Baspa 103 River, a stream of the Sutlej River and finally joins the Indus River system. The study area has 104 been presented in Figure 1, where locations of the instruments like the Automatic Weather 105 106 Station (AWS) discharge gauge have also been shown. The watershed and glaciated area in the catchment have also been presented along with the contour map. This catchment receives 107 precipitation from the winter westerlies and the monsoon in summer (Singh et al. 2018), but 108 109 summer precipitation dominates winter precipitation. The winter precipitation in this region is received through the Western Disturbance (WD) (Dimri 2004). This region experiences 110 ablation from May to September (Kumar et al., 2016). It covers an area of 38.13 km² above the 111 discharge site. The hypsometric distribution of the catchment is presented in Figure 2. 112

Hypsometry distribution clearly shows that 75% of the catchment has a non-glaciated area 113 while 25% is glaciated. In the glaciated area, the debris-covered glacier has its share of 5%, 114 while the debris-free glacier has 20%. The rocks of this region resemble the Higher Himalayan 115 Crystalline. It contains pelitic and psammopelitic meta-sediments having acid and basic 116 intrusive. Granite and gneiss rocks in the Himalayan region have a familiar presence of late-117 stage pegmatitic veins feldspar (Kumar et al., 1987). Rohtang gneiss is the principal constituent 118 119 of the Himalayan glaciers. Chalcopyrites' presence is also noticed in lateral morainic deposits of the Himalayas. The morphology of the Shaune Garang catchment describing the area vs 120 121 aspect and area vs slope has been shown in Figure 3. The aspect ratio suggests that most of the 122 part of the catchment falls in the East and South, Southwest and West direction and receives

higher solar insolation, making it prone to weathering. Further, the primary area falls in the
middle slope zone of 24° to 40° and the higher slope zone 40° to 56°, which transfers the
weathered materials very quickly to the glacier.

126 Sampling and analysis protocol

In the higher Himalayas, the physical accessibility is limited to the summer season. Water 127 128 samplings were performed during the summer seasons of 2016 and 2017 at the selected locations of the channel (Figure 1). Selected cations and anions of glacial meltwater were 129 examined in 2016 and 2017. Fifty-three water samples in 2016 and 26 water samples in 2017 130 were collected during ablation season. The sampling sites were chosen as per the geology, 131 altitude, terrain, and convergence of tributaries. Polyethylene sample bottles (250 ml) were 132 rinsed with distilled water after being cleaned with nitric acid for accuracy in data. Water 133 samples were taken from 20 cm deep to have a well-mixed concentration. Sampled glacial 134 135 meltwater was filtered on 0.45 µm Whatman filter paper. The conductivity and pH measurements have been performed in the field by a handheld multi-parameter instrument 136 (HANNA, model No. HI9829). The essential cations (Ca²⁺, Mg²⁺, K⁺ and Na⁺) analyses were 137 performed using Atomic Absorption Spectroscopy. The instrument has a precession of 0.05 138 parts per million (ppm) for Ca²⁺, Mg²⁺, K⁺, Na⁺, and 0.01 parts per million (ppm) for the 139 remaining parameters. Ion chromatography was used to analyze the anions (Cl^{-} , SO_4^{2-} and 140 NO₃⁻). The bicarbonate concentration was based on the results of the titration method (APHA 141 2005) and the charge balance method. The precautions were made as per the specified norms, 142 143 and we used non-powder vinyl cleanroom gloves and masks for sampling and analysis. A new standard of identified concentration and procedural blank was analyzed for every analytical 144 run. During the investigation, no detectable contamination was obtained. The data from physio-145 chemical analysis of the meltwater samples were subjected to multivariate statistical analysis. 146

147 The study used multivariate statistical tests to examine relationships among multiple variables

in the data set. Excel add-on XLSTAT was used for the analysis of normalized data under PCA.
Bartlett's sphericity test was applied to both years' melting season data, and a correlation matrix
was prepared. The principal component analysis has been analyzed to understand problems
under different measurement scales of the original variable, which is avoided by diagonalizing
the correlation matrix. The dissolved ions composition of the meltwater from Shaune Garang
glacier were subjected for the error analysis in the charge balance, which has been computed
through the given formula:

155
$$CBE = \frac{(TZ^{+} - TZ^{-})}{(TZ^{+} + TZ^{-})} \times 100$$

156

Equation 1

157 Where, (TZ^+) = total cations and (TZ^-) = total anions

The errors of TZ^+ and TZ^- were < 10 % for two consecutive melting periods (2016 and 2017), which is indicative of the good quality of data.

160 Remote sensing data and characteristics

Air temperature of glacier surface was measured in ablation season (June to October) during 161 the study period. Satellite data has been validated by the ground-based temperature records of 162 snow, ice, and debris-covered glacier at twenty sites during the satellite passes at 10:30 AM 163 164 over the study area. ASTER data were used in this investigation to obtain lithological and mineralogical evidence from the Shaune Garang catchment. A subset of the high-resolution 165 and cloud-free images (ASTER) were used throughout the study to determine the precise 166 lithological mapping of the exposed rock present in the catchment. The imagery of nine bands 167 was used in this study, with band-1 being blue (0.43 - 0.45 µm), band-2 being blue (0.45 -168 0.51µm), band-3 being green (0.53 - 0.59 µm), band-4 being red (0.64 - 0.67 µm), band-5 being 169

170 near-infrared (0.85 - 0.88 μ m), band-6 being shortwave infrared (1.57 - 1.65 m), band-7: 171 shortwave infrared (2.11 - 2.29 μ m), band-8: panchromatic (0.50 - 0.68 μ m), and band-9: cirus 172 (1.36 - 1.38 μ m) were used. Respective bandwidths and subsystems characteristics are 173 presented in Table1.

174 **Result and discussion**

175 Hydrogeochemistry of glacial meltwater

The physical analysis of meltwater indicates a little alkaline with pH ranging from 6.86-8.56 176 with an average value of 7.45 ± 0.48 and 7.45 ± 0.45 for 2016 and 2017, respectively (Table 2). 177 Higher pH indicates that the process of dissolution is higher due to the more considerable 178 contact period with rock, soil and rainwater. These would have imparted alkalinity to the 179 meltwater (Kumar et al. 2014). Electrical conductivity indirectly measures the mineralization 180 that explains the ionic strength of water (Kumar et al., 2019). The standard value of electrical 181 182 conductivity was 86.14 \pm 16.96 μ S/cm in 2016 and 91.25 \pm 16.62 μ S/cm in 2017. The higher EC must result from the weathering, evaporation, and crystallization processes. Furthermore, 183 the lesser conductivity in glacier discharge is influenced by increased precipitation making 184 higher discharge and decreased influence of the evaporite dissolution process. The measured 185 value of EC suggests that hydrochemistry of location is regulated through the interface of water 186 and rock and depends on the weathering of rocks. Different rocks and their solubility influence 187 the proportional concentration of ions in glacial meltwater (Pant et al. 2021). Table 2 displays 188 the distribution of dissolved ionic concentrations with standard deviation in the meltwater 189 190 discharge of the Shaune Garang glacier.

The results presented in Table 2 indicate that Ca^{2+} contributes 39.57% and 42.53% in the total cationic budget in both the consecutive study periods 2016 and 2017. However, $Ca^{2+} + Mg^{2+}$ contributes 82.10% and 71.02% of the total cationic budget in the catchment. The other two

cations, Na⁺ and K⁺, contribute only 16.78 % and 17.57 %, respectively, during the study period 194 2016 and 2017. Bicarbonate (HCO₃⁻) is the most dominant anion contributing 62.18 % and 195 54.44 %, respectively, in the total anionic budget of the ablation period of 2016 and 2017. Its 196 average concentration was observed as $369.65 \pm 79.41 \ \mu eq/l$, and $316.73 \pm 83.23 \ \mu eq/l$ in the 197 consecutive study period. Sulphate (31.10% and 38.11%) was the second most dominant 198 anion, followed by chloride (5.58% and 6.64%) and Nitrate (1.18% and 0.79%) during 199 200 consecutive years' observation. The dominance of bicarbonate in Shaune Garang Catchment is due to the silicate dominating geomorphology of the catchment. According to the findings, 201 202 weathering of silicate minerals is less visible than carbonate minerals.

Figure 4 displays the concentration of different anions and cations and electrical conductivity 203 for the glacier's meltwater at different parts of the Indian Himalaya. The concentration of 204 cations and anions varies as per the morphology of rocks in the catchments and weather system. 205 Meltwater draining from the Himalayan region shows the dominance of Ca²⁺ and HCO₃⁻, 206 whereas Bagni, Chaturangi, Gangotri and Dudu glaciers demonstrate the dominance of SO₄²⁻ 207 208 in their catchment. The dominating presence of silicate-bearing rocks is the important factor for the higher concentration of bicarbonate (HCO₃⁻) in the meltwater of the Himalayan glacier. 209 In addition, the dominancy of SO_4^{2-} in the Bagni, Chaturangi, Gangotri and Dudu glaciers could 210 be due to pyrite oxidation that enhances sulphate concentration. Cl⁻ and SO₄²⁻ domination is 211 influenced by halite and sulfide oxidation and weathering of soft sulphate minerals such as 212 gypsum (Thomas et al., 2015). The chemical composition analysis reflects the dominance of 213 bicarbonate (HCO₃⁻) as an anion in most of the glacial meltwater in the Himalayan region due 214 215 to the dissolution of atmospheric carbon dioxide and carbonate (Sharma et al., 2013; Kumar et al., 2014; Singh et al., 2015). Concentration of cations in the meltwater of Dokariani, Bara 216 Shigri and Gangotri glaciers follows a trend like $Ca^{2+} > Mg^{2+} > K^+ > Na^+$ while it follows a 217 trend of $Ca^{2+} > Mg^{2+} > Na^+ > K^+$ for meltwater of Kafni and Chhota Shigri, the only alteration 218

in the concentration of Na⁺ and K⁺ for different basins. However, the Bagni glacier meltwater 219 showed the Potassium ion as the second most abundant. In the Bara Shigri glacial, meltwater 220 concentration of cations varied as $Ca^{2+} > Mg^{2+} > Na^+ > K^+$ like the Chhota Shigri of its vicinity 221 and the Kafni of Kumanyu Himalaya whereas anions concentration followed the pattern of 222 $HCO_3^- > SO_4^{2-} > NO_3^-$. It has been observed from the comparative analysis in Figure 4 that the 223 central Indian Himalayan glacier's meltwater has the highest concentration of Ca²⁺ cation, and 224 225 a similar observation is from the present study. Interestingly, the graph shows a higher concentration of anions and cations in discharge from glaciers located in the central part of the 226 227 Indian Himalayan region than the glaciers in the western part.

228 Hydro-geochemical process in the glacial catchment

Dissolved ions in the glacial meltwater are generally contributed through rock weathering, 229 precipitation, anthropogenic influence, and atmospheric conditions (Jeelani 2011; Kumar et al. 230 2019). Generally, the chemical composition of glacial meltwater is governed by the chemical 231 weathering between the interaction of water and bedrock beneath the glacier (Kumar et al., 232 2009; Kumar et al., 2014; Singh et al., 2017). The cation and anion in the glacier discharge are 233 234 elucidated concerning the nature of rock and its weathering processes. Dissolved solute particles present in the glacial melt are determined by the processes involved in the glacial 235 236 environment. The interrelationship between the physical and chemical parameters are presented through a scatter plot (Figure 5) of $(Ca^{2+}+Mg^{2+})$ and $(Na^{+}+K^{+})$ against total cation 237 (TZ⁺). A positive correlation is observed between ($Ca^{2+}+Mg^{2+}$) and TZ⁺. It further shows a 238 ratio ranging from 0.75 to 0.85 with an average equivalent value of 0.75 ± 0.05 during the study 239 period. 240

The result demonstrates that the impact of $Ca^{2+}+Mg^{2+}$ in the glacial meltwater is comparatively high compared to the total cation TZ⁺. The ratio of Ca^{2+} and Mg^{2+} determines the input source

of calcium and magnesium ions in water. $Ca^{2+}/Mg^{2+} \le 1$ (Table 3) indicates a process of 243 dolomite dissolution, and a value >1 recommends the dominance of silicate weathering in water 244 (Kumar and Singh 2015). As a result, silicate weathering could be a factor in the dominant 245 concentration of Ca^{2+} and Mg^{2+} among cations in the Shaune Garang glacial discharge. The 246 scatter plot (Figure 5) among Na⁺+K⁺ and TZ⁺ displays a small contribution of Na⁺+K⁺ in total 247 dissolved ion with 0.24 \pm 0.04 and 0.25 \pm 0.08 during both ablation years. The results reveal 248 249 carbonate weathering as a leading factor in the glacial meltwater ionic characteristics of the Shaune Garang catchment. The large equivalent ratios 3.23 ±0.75 and 3.28 ±1.14 for 250 $(Ca^{2+}+Mg^{2+})/(Na^{+}+K^{+})$ in the consecutive melting period of 2016 and 2017 further strengthen 251 the understanding of carbonate weathering dominance in the catchment. High ratio of 252 $(Ca^{2+}+Mg^{2+})/(Na^{+}+K^{+})$ and $(Ca^{2+}+Mg^{2+})/TZ^{+}$ in the glacial meltwater demonstrate that hydro-253 geochemistry of the meltwater of Shaune Garang catchment is mainly administrated by CO₃ 254 weathering with a minor contribution of SiO₂. However, the evaporation process enhances the 255 TDS concentration in water (Prasanna et al. 2010; Xing et al. 2013). The average ratio of 256 Na^{+}/Cl^{-} was measured to be 4.77 ±2.27 and 3.83 ±1.90 in 2016 and 2017 (Table 3). The Na^{+}/Cl^{-} 257 ratio indicates a minor contribution of atmospheric constituents in the chemical 258 characterization of meltwater of the catchment. 259

The ion exchange process in the water is mainly defined because of $(Ca^{2+}+Mg^{2+})$ versus 260 $(HCO_3^- + SO_4^{2-})$ (Srinivasamoorthy et al., 2008). Dominant dissolution process of calcite, 261 dolomite, and gypsum ion exchange may shift the points rightward owing to excess of (HCO₃⁻ 262 +SO 4^{2-}). Further, the reverse ion exchange process turns leftward due to a surplus (Ca²⁺+Mg²⁺). 263 $(Ca^{2+}+Mg^{2+})$ versus $(HCO_3^- +SO_4^{2-})$ indicates carbonate and silicate weathering with ion 264 exchange as leading geochemical processes in the catchment (Figure 6). The diagram displays 265 that contribution of $(HCO_3^- + SO_4^{2^-})$ to the total ionic concentration in the glacial meltwater is 266 greater than the ($Ca^{2+}+Mg^{2+}$), indicating an excess of ($HCO_3^-+SO_4^{2-}$) which is contributed by 267

silicate weathering. The dominance of calcium and magnesium ions is calculated through 268 Ca^{2+}/Na^{+} and Mg^{2+}/Na^{+} ratios, a product of weathering of CO₃ and SiO₂. The Ca^{2+}/Na^{+} ratio 269 was observed as 2.57 \pm 0.80 and 2.64 \pm 1.06, while Mg²⁺/Na⁺ was respectively 2.16 \pm 0.67 and 270 2.07 ±0.81 in the meltwater of Shaune Garang glacier (Table 3). This ratio shows the 271 dominance of Ca²⁺ and Mg²⁺ over Na⁺. The result further confirms that hydro-geochemistry is 272 governed by weathering of CO₃ minerals in the catchment. Hydrogen ion availability is 273 responsible for rapid CO₃ weathering (Das and Kaur 2001). Na⁺ normalizes Ca²⁺ and HCO₃⁻ 274 concentration and determines the effect of SiO₂ weathering, evaporative dissolution or CO₃ 275 276 weathering in meltwater (Kumar et al., 2015). To understand the chemical weathering, sulphate mass fraction (SMF), and the ratio of sulphate (SO_4^{2-}) to $(SO_4^{2-} + HCO_3^{-})$ have been calculated 277 in the study catchment. The chemical characteristics of meltwater show the importance of 278 carbonation if the SMF value is (<0.5). The SMF value indicates the chemical attributes of 279 meltwater affected by sulfide oxidation and the termination of CO₃ (Tranter et al. 1993). In the 280 Shaune Garang catchment, an average SMF value of 0.33 ± 0.07 and 0.41 ± 0.09 , respectively, 281 during the study period 2016 and 2017 indicates the dissolution of carbonate and sulfide 282 oxidation. In Addition, C-ratio ($HCO_3^{-}/HCO_3^{-}+SO_4^{2-}$) has also been calculated to find the 283 significance of proton-producing effects necessary for the chemical weathering of carbonate 284 rocks. During 2016 and 2017, the C-ratios were 0.67 \pm 0.07 and 0.59 \pm 0.09, demonstrating the 285 domination of the carbonate and sulphate weathering processes. 286

287 Mineral mapping

The Short Wavelength Infra-Red (SWIR) and Thermal Infrared (TIR) spectral resolution agree for mapping surface mineralogy. These spectral bands are available in (ASTER) and have been used to map the distribution of minerals on supraglacial debris. Minerals are mapped through the band indices like "SWIR indices", "TIR indices", and "TIR emissivity silica weight per cent" in the Shaune Garang catchment. The mineral measurement reflects the primary presence of quartz, feldspar, carbonate, and mica. High altitude glacier debris reflected the fact of "quartz, feldspar as calcium albite, and mica as biotite". The debris on the Shaune Garang glacier is dominated by muscovite (mica), calcium albite (feldspar) and quartz. Though in lesser quantity, the presence of calcite has also been noticed. To create thematic mineral abundance maps and quantitative estimation of minerals, "SWIR and TIR indices" have also been used (Ninomiya, 2004).

299 SWIR indices

300 Short Wavelength Infrared (SWIR) mineral indices were used to wavelength-dependent 301 absorption patterns in estimating minerals in the catchment. The SWIR mineral indices were 302 used to evaluate the mineral's dominance in the catchment. Equations 1, 2, 3 and 4 have been 303 used respectively for understanding the dominance of layered silicate (LS), calcite (CA), 304 hydroxyl-bearing (OH), and Alunite (AL).

305	$\mathbf{LS} = \frac{(\mathrm{AST4} \times \mathrm{AST8})}{(\mathrm{AST5} \times \mathrm{AST6})}$		
306	Equation		1
310	$\mathbf{CA} = \frac{(\text{AST6} \times \text{AST9})}{\text{AST8}^2}$		
307		Equation	2
311	$\mathbf{OH} = \frac{(\text{AST4} \times \text{AST7})}{\text{AST6}^2}$		
308		Equation	3
312	$\mathbf{AL} = \frac{\mathbf{AST7}^2}{(\mathbf{AST5} \times \mathbf{AST8})}$		
309		Equation	4

313 Where,

The varying indices are related to the variable absorption properties, which helps measure the 315 316 types of minerals. A sensor, "radiance band ratios", can reduce the influence of the atmosphere and the topography of a region and the variation in illuminance (Abrams et al., 1983; Mather, 317 318 1987). The evidence also indicates the nonsignificant evidence in "single band or three-band true or false colour composite imageries". It is also helpful in having the quantitative estimation 319 of mineral abundances. In this study, images of the 4-shortwave infrared (SWIR) mineral 320 indices are displayed in Figure 7, reflecting the relative dominance of minerals and their 321 presence on the surface. Alunite has been most dominant and abundant in higher altitudes up 322 to the accumulation zone. "Layered silicates" and "hydroxyl-bearing minerals" are less 323 productive, while "calcite and hydroxyl-bearing minerals" varies location-wise. Alunite index 324 displays little white patches in the higher region with high abundance. Figure 7 shows 325 kinematics and pulse flow movements of layered silicate debris, which can be understood 326 327 through their variability and abundance. The evidence of Alunite at a higher altitude might be due to its formation mechanism. The formation of Alunite through the reaction of sulfuric acids 328 with potassium-rich feldspars is called "alunitization". Layered silicates and "hydroxyl-bearing 329 minerals" are in short supply, but "calcite and hydroxyl-bearing" minerals vary significantly 330 within the catchment. Layered silicate consists of octahedral layers bound to the tetrahedral 331 and primary component of soil. Its distribution within the catchment at lower altitudes implies 332 the weathering mechanism due to meltwater and parental rock interaction. They have been the 333 excellent water trapping mechanism held between layers. The essential minerals in layered 334 335 silicates are kaolinite, nacrite and dickite.

336 **TIR indices**

To evaluate various minerals in the Shaune Garang catchment area, Thermal infrared (TIR) 337 mineral indices of Carbonate, Quartz and Mafic were used. The thermal spectrum is 338 instrumental in distinguishing the geology of earthy minerals, where TIR satellite spatial 339 resolution is noticeably lesser than VNIR or SWIR (VNIR 15 m, SWIR 30 m, TIR 90 m). 340 However, TIR is exclusive in targeting the profusion of carbonate, quartz, and silicate minerals. 341 Band ratios derived from TIR estimate carbonate, quartz and silica bearing lithology (Figure 342 343 8). Equations 5, 6 and 7 were used for Carbonate Index (CI), Quartz Index (QI), and Mafic Index (MI), respectively. 344



359 of dolomite minerals. Minerals of carbonates with "hydrothermal origin" are very challenging

to identify through a CI map due to an inadequate percentage of carbonate presence. However, 360 calcite-bearing propylitic alteration is accredited as the "ASTER TIR" feature. The chlorite and 361 epidote had lower emissivity between TIR bands 11 and 13 but slightly higher between 13 and 362 14 (Salisbury et al. 1992). Aspects of the spectrum with these characteristics resemble the mafic 363 index minerals. Mafic and quartz index minerals found uneven distribution within the 364 catchment, but carbonate minerals were found at lower altitudes along the riverside. The reason 365 366 behind the occurrence of carbonate minerals along river channels might be due to the higher weathering across the river. 367

368 Chemometric Analysis

369 Principal component and factor analysis

Excel add-on XLSTAT was used for the analysis of normalized data under PCA. The sphericity test of Bartlett was performed on the data of both years. The Bartlett sphericity test shows that observed χ^2 (342.85) is considerably more significant than the critical χ^2 (85.96) in 2016 and χ^2 (observed) = 125.25 larger than the critical value χ^2 (critical) = 48.3 in 2017. The principal component analysis helped to understand problems under different measurement scales of the original variable avoided by diagonalizing the correlation matrix.

Table 4 demonstrates the PC value of more than 1, which explains 72.1 % of the total variance 376 of four PCs. PCs 1, 2, 3 and 4 are capable of explaining 39.21%, 12.91%, 10.24% and 9.74% 377 of variance in 2016. Similarly, in 2017 scree plot (Figure 9) shows four PCs, which explains 378 69.9.1% of the total variance. PC 1, 2, 3 and 4 can explain 26.62 %, 20.12 %, 12.64% and 379 10.52 % of variance. Table 4 further depicts the Eigenvalues, the percentage of variance 380 calculated through varimax rotation matrix with Kaiser Normalization and rotated factor, and 381 the percentage of variance in each PC. High SO_4^{2-} and K⁺ loadings are observed in both the 382 years, indicating silicate weathering dominance in the catchment. Moderately high loading 383

values of Ca^{2+} , Mg^{2+} and Na^{+} in both years indicate the dominance of the process, which is prevailing in factor 1.

Table 5 indicates that factor 4 shows the negative pH in both the years and the acceptable value of Cl⁻ in the consecutive study period. The first two principal component loading are presented to understand the grouping and relationship of all chemical parameters. The PC loading has been calculated to understand correlations among variables and know the most influential variables. It could result from the minerals present in the soil (Yakubo et al., 2009). Throughout the study, higher loadings in Na⁺ and Mg²⁺ may be accredited to the ionic conversation between water through dissolution minerals containing sodium.

Statistically, the coefficient of determination (R^2) indicates one variable's level of statistical 393 agreement with another. Here, it is applied among the hydrogeochemical parameters and 394 represented in (Table 6) during the study period. Water chemistry parameters such as EC and 395 Na⁺ are highly interrelated with Ca²⁺, Mg²⁺, and HCO₃⁻. Similarly, a decent relationship among 396 $(Ca^{2+}-Mg^{2+})$, $(Ca^{2+}-HCO_3^{-})$ and $(Mg^{2+}-HCO_3^{-})$ has been observed. The above parameters 397 have a good positive correlation ($R^2 > 0.5$) and are an indicator of control by these parameters 398 in the solute chemistry of the study region. The strong correlation between parameters such as 399 Ca^{2+} , Mg^{2+} , $(Ca^{2+} - HCO_3^{-})$, $(Mg^{2+} - HCO_3^{-})$ indicates strong carbonate weathering (Singh et 400 al., 2017). In the case of sulphate (SO_4^{2-}) ion concentration, it shows a good relationship with 401 Ca^{2+} and Mg^{2+} , indicating the sulphate mineral's high dissolution in the glacier's catchment. 402

403 Hydro-geochemical facies of the glacial meltwater

Hydrogeochemical facies of the meltwater helps interpret the dominant anions and cations,
which have been determined through a Piper plot (Figure 10). It is used to find similarities and
dissimilarities among all water types, where the analogous water qualities fall together (Todd,
2001). In the cation plot, it can be seen that most of the water is concentrated in a trilinear

pattern in the middle, indicating that it is a mixed water type. The calcium ions predominate in 408 the glacial discharge. The hydro-geochemical cations in the bottom left triangle of Shaune 409 Garang glacial discharge prove calcium ions' dominance. It substantiates the conclusions 410 reached in the sections on hydro-geochemistry and hydro-geochemical processes, both 411 included in the previous section. Slightly elevated sodium and potassium ion concentrations in 412 several samples confirm the presence of sources at the various sampling locations of the Shaune 413 414 Garang catchment. This must be contributing to the overall cation concentration (Karim and Veizer, 2000; Ravikumar, 2017). The Piper plot aids in the understanding of the fact, Ca^{2+} , 415 Mg^{2+} and HCO_3^{-} are the most prevalent ions in the Shaune Garang catchment. The average 416 percentage value of $(Ca^{2+}+Mg^{2+})$ is about 77% and 81%, respectively in the years 2016 and 417 2017; however, for the $(Na^+ + K^+)$ it showed about 33 % and 29%, respectively, demonstrating 418 that alkaline earth metals are prevailing over alkali metals (Singh et al., 2017). It further 419 supports the dominance of dolomitic limestone containing gypsum and pyrite in the region 420 (Figures 7 and 8). The piper diagram demonstrates that carbonate type weathering has been 421 more instrumental in governing hydro-geochemistry in this catchment. The figure indicates the 422 presence of the $(Ca^{2+} - HCO_3^{-})$ type of water with little influence from $(Ca^{2+} - SO_4^{2-})$ type. 423 Significant ions and TDS were found in higher concentrations in this catchment, indicating 424 more significant interactive processes between rock materials and water having influential 425 weathering due to moisture. 426

427 Apart from the Piper plot, Gibb's diagram (Gibbs 1970) was applied to understand how hydro-428 geochemical techniques such as precipitation, rock-water interface, and vaporization impact 429 the environment's hydrogeology. According to the Gibbs diagram (Figure 11), the chemical 430 weathering of rock minerals and a minimal extent of evaporation crystallisation are the 431 essential variables to consider the meltwater quality in the Shaune Garang catchment. Gibb's 432 diagram advocates the higher rock-water interaction resulting in higher ionic concentration in meltwater. Chemical weathering, carbonate dissolution and ionic exchange between water and
clay indicate the rock-water interaction processes (Kumar et al., 2014). Increased evaporation,
chemical weathering and anthropogenetic actions raise the total dissolved solids. Furthermore,
the findings show that water contamination from poor sanitation has increased Na⁺ and Cl⁻ ions
and increased total dissolved solids (TDS) (Kumar et al., 2014).

438 Conclusion

The current study focused on the hydro-chemical analysis of meltwater from the Shaune 439 Garang catchment, located within the Baspa Basin. According to the findings, meltwater is 440 slightly alkaline with Ca²⁺ and HCO₃⁻ the most prevalent ions during the study. Ca²⁺ and Mg²⁺ 441 were the dominant cations constituting (41.03%, 42.53%) and (34.84%, 32.89%) of the total 442 cationic budget in the consecutive study period. The predominance of carbonate weathering is 443 indicated by the ratio of $(Ca^{2+}+Mg^{2+})/(Na^{+}+K^{+})$ and $(Ca^{2+}+Mg^{2+})/(TZ^{+})$ and a strong positive 444 association between Ca^{2+} – Mg^{2+} , Ca^{2+} – HCO_3^- and Mg^{2+} – HCO_3^- . Piper plot demonstrated that 445 alkaline earth metal dominates alkali metal, and weak acid exceeds strong acid. This plot also 446 indicated that the Ca^{2+} -HCO₃⁻ is most influential, assessed by Mg²⁺-HCO₃⁻ type of water in 447 this area. The Gibbs plot also revealed that rock corrosion is the primary process regulating 448 meltwater concentration. The values in the chloro-alkaline indices in this study were negative, 449 indicating the conversation of Ca²⁺ and Mg²⁺ ions by Na⁺ and K⁺ ions of rock material. The 450 geological mapping of the catchment has been done on varying scales, including diversified 451 rocks and unconsolidated surface materials that possess "quartz and carbonate minerals". The 452 results were verified through the geological, stratigraphic, and structural maps in the 453 multifaceted lithological terrain of the region. Based on the lithological map of the Shaune 454 455 Garang catchment, Layered silicates and "hydroxyl-bearing minerals" are less abundant. However, "calcite and hydroxyl-bearing minerals" have significantly varying availability in the 456 catchment. The distribution of Layered silicate minerals within the catchment is majorly found 457

at lower altitudes, which implies the weathering mechanism due to the interaction of meltwater 458 and parental rock. The chemometric analysis includes a principal component (PC), 459 eigenvalues, the percentage of variance calculated through varimax rotation matrix with Kaiser 460 Normalization and rotated factor, and the percentage of variance in each PC. High SO_4^{2-} and 461 K⁺ loadings are observed in both years, indicating silicate weathering dominance in the 462 catchment. Moderately high loading values of Ca^{2+} , Mg^{2+} and Na^{+} in both years indicate the 463 dominance of the process, which is prevailing in factor 1. The strong correlation between 464 parameters such as Ca²⁺, Mg²⁺, (Ca²⁺ - HCO₃⁻), (Mg²⁺ - HCO₃⁻) indicates intense carbonate 465 weathering (Singh et al. 2017). In the case of sulphate (SO_4^{2-}) ion concentration, it shows a 466 good relationship with Ca^{2+} and Mg^{2+} , indicating the sulphate mineral's high dissolution in the 467 glacier's catchment. 468

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476 **Declaration of interests**

477 The authors declare no competing financial or personal interests that influenced their work.

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704	<u>List of figures</u>
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Figure 8: Mineralogy maps based on the "ASTER TIR" of Shaune Garang catchment's radiance properties. The indexes are "Carbonate Index (CI), Quartz Index (QI), and Mafic Index (MI)".

- **Figure 9:** Scree plot graph between eigenvalue and factors for 2016 and 2017.

Figure 10: Piper plot for concentration of ions during study period 2016 and 2017 in the Shaune Garang catchment.

Figure 11: Gibb's diagram represents controlling factors of meltwater quality.



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Sampling locations are numbered in ascending order from higher altitudes to lower altitudes.

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Figure 4: The average hydro-geochemical characteristics of glacial meltwater from Shaune





Figure 5: Scatter plot of $(Ca^{2+}+Mg^{2+})$ against TZ⁺ and (Na^++K^+) against total cation (TZ⁺) during the study period of 2016 and 2017 from Shaune Garang glacier catchment.



Figure 6: Scatter plot of $(Ca^{2+}+Mg^{2+})$ against $(HCO_3^-+SO_4^{2-})$ and $(Ca^{2+}+Mg^{2+})$ against SO_4^{2-} during the study period of 2016 and 2017 from Shaune Garang glacier catchment.



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Figure 7: Mineralogical maps based on the radiance properties of ASTER SWIR of Shaune
Garang catchment. The indexes are represented through LS, CA, AL, and OH for layered
silicate, calcite, Alunite, and hydroxyl bearing materials.



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Figure 8: Mineralogy maps based on the "ASTER TIR" of Shaune Garang catchment's
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772 Shaune Garang catchment.



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