#### **ORIGINAL PAPER**



# **Assessment of water recharge source of geothermal systems in Garhwal Himalaya (India)**

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#### **Abstract**

The geothermal systems in the Himalaya are complex, and their genesis, circulation pattern and processes of sustenance are largely unknown. The present study aims to systematically analyse the characteristics of oxygen and hydrogen isotopes, major ion data and strontium ratio ( ${}^{87}Sr/{}^{86}Sr$ ) of geothermal systems, river waters and rainwater to understand the movement of groundwater and the mechanisms for the formation of geothermal systems in the region. Further, the strontium isotope was also used to understand the fuid source of geothermal systems. Field observations show that the geothermal waters have a higher temperature, hydraulic pressure and elevated  $\delta D$  and  $\delta^{18}O$  values than river waters. Thus, large river systems are not the principal recharge source of geothermal systems. Meteoric water (rain and snowmelt) in high mountains can infltrate and circulate deep down the active tectonic belts or sutures and recharge geothermal systems. The cold surface water evolves into high-temperature thermal water after deep circulation and is discharged as a geothermal spring at the surface, under a high water-head diference. Therefore, the large-scale geothermal systems in the Garhwal Himalaya develop and are maintained by rapid groundwater circulation and interaction with a heat source. Further, the water temperatures of these systems in Garhwal Himalaya have remained the same over the period 1975–1994 (Geological Survey of India) and 2010–2016 (this study) with an error of  $< 5\%$ .

**Keywords** Geothermal systems · Groundwater · Isotopes · Himalaya

# **Introduction**

The Himalayan range is an active mountain chain that hosts many geothermal systems in natural and artesian conditions that eject hot waters and volatiles with varied temperatures and chemical compositions. The Himalaya-Karakoram region stretching from Pamir to Yunnan has documented ~ 600 geothermal systems (Gyanprakash and Raina [1975;](#page-16-0) Tong and Zhang [1981\)](#page-17-0). The South Asia region is under active tectonic process over the last 50 million years, producing the Himalayan orogeny by continent–continent collision. The underthrusting of the Indian plate beneath the Eurasian plate has led to the rise of the Himalaya through the process of crustal thickening

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 $\boxtimes$  Akshava Verma akshayaverma5@gmail.com (Molnar et al. [1987](#page-16-1); Bilham et al. [1997;](#page-15-0) Hochstein and Regenauer [1998](#page-16-2)) and crustal melting (Searle et al. [1997](#page-16-3); Henry et al. [1997](#page-16-4)). Steep geothermal gradients that can force hydrothermal circulations along with degassing of volatiles are associated with the rapid uplift and exhumation in such mountain belts (Koons and Craw [1991;](#page-16-5) Jenkin et al. [1994;](#page-16-6) Koons et al. [1998;](#page-16-7) Tiwari et al. [2020\)](#page-17-1). Therefore, geothermal circulation can be an important phenomenon in the thermal budget of an orogeny (Ingebritsen et al. [2001;](#page-16-8) Forster and Smith [1989;](#page-15-1) Manga [1998](#page-16-9); Derry et al. [2009\)](#page-15-2). The world's rarest and distinct geothermal systems have developed in the deeply incised valleys beside the main boundary thrust (MBT), the main central thrust (MCT) and the Indus Tsangpo suture zone (ITSZ) with the upliftment of the Himalayan mountain chain (Evans [2003](#page-15-3); Evans et al. [2004](#page-15-4); Becker et al. [2008\)](#page-15-5). The distribution of geothermal systems is asymmetric to the Himalayan axis. However, several of these systems in northwest Himalaya, especially those with surface temperatures of  $T_s \sim 90 \degree C$ , fall next to the main thrust zones, indicating that these have structural control (Hochstein and Regenauer [1998](#page-16-2);

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Newell et al. [2008;](#page-16-10) Evans et al. [2008;](#page-15-6) Rai et al. [2015](#page-16-11); Tiwari et al. [2016b](#page-17-2)).

Geothermal systems are of cyclic and storage types globally. Himalayan geothermal systems fall in the cyclic category, where water is meteoric (Gupta and Roy [2006](#page-16-12)), percolates to deeper depths and then heats and upwells to the surface (Tiwari [2014\)](#page-17-3). Some previous studies have focused on the formation of geothermal systems in the Himalayan region and assume that meteoric waters infltrate and circulate within a restricted space where heat is transferred from a source to the sink to recharge the geothermal systems (Chandrajith et al. [2012](#page-15-7); Tan et al. [2014](#page-17-4); Tiwari et al. [2016a\)](#page-17-5).

Based on geophysical exploration, it has been confrmed that the magmatic input through crustal remelting is the principal heat source in the Tibetan Plateau (Tan et al. [2014](#page-17-4)). The heat source for geothermal systems could be a lowtemperature system, in which temperature increases with depth or a magmatic intrusion  $(>600 \degree C)$  that has reached relatively shallow depths of 2 to 5 km (Harinarayana et al. [2006\)](#page-16-13). The circulating fuid reservoir could extract heat from a volume of hot permeable rocks overlain by an impermeable layer. The geothermal fuid is water, which exists as a liquid or vapour phase depending on its temperature and pressure. This water often carries chemical constituents like boron (B), lithium (Li), caesium (Cs), rubidium (Rb), strontium (Sr), arsenic (As), iron (Fe) and uranium (U) (Gupta and Agarwal [2001;](#page-15-8) Evans et al. 2003; Tan et al. [2014](#page-17-4)) and gases such as  $CO_2$ ,  $H_2S$ ,  $SO_2$ , He and  $CH_4$  (Cinti et al. [2009](#page-15-9)). Zones of geomagnetic anomalies, shallow earthquakes and gravity anomalies are generally associated with these noneruptive systems (Tiwari et al. [2020](#page-17-1)) and other processes in these settings like heat fow and migration of magmatic fuids from the deep to the surface (Azeez and Harinarayana [2007](#page-15-10); Veeraswamy et al. [2010;](#page-17-6) Rawat [2012](#page-16-14)). However, previous studies have discussed the heat source of geothermal systems based on geological and geophysical data. Only a few have used stable water isotopes to observe and characterise the relationship between the formation mechanism, heat source and geochemistry of geothermal waters (Chandrajith et al. [2012](#page-15-7); Tan et al. [2014](#page-17-4); Tiwari et al. [2016a](#page-17-5)).

Given the diverse stable water isotopic compositions of geothermal springs, cold springs, river water, rainfall, snowfall and glacial meltwater, it offers a tremendous opportunity to trace the groundwater circulation, including recharge, runoff and discharge (Clayton et al. [1966;](#page-15-11) Clark and Fritz [1997](#page-15-12); Tan et al. [2012](#page-17-7), [2014](#page-17-4); Tiwari et al. [2016a](#page-17-5)). This work aims to scientifcally analyse the characteristics of stable water (hydrogen and oxygen) and radiogenic strontium isotopes (87Sr/86Sr) in diferent components of the hydrological cycle (geothermal systems, cold springs, river water, rainfall, snowfall and glacial meltwater) and to understand the circulation and source of water, heat source and hydraulic

gradients to maintain the geothermal systems in Garhwal Himalaya.

#### **Study area**

The Garhwal region of the central Himalaya receives both the Indian summer monsoon (ISM) and westerlies which infuence the rivers and streams draining the area. The Ganga River system has two major proglacial rivers, i.e. Bhagirathi and Alaknanda, and spreads over an area of  $\sim$  30,000 km<sup>2</sup>. Major streams of the region forming the headwaters of the Ganga carry 70–80% of their annual flow during the summer monsoonal months (Bruijnzeel and Bremmer [1989\)](#page-15-13). Most of the rainfall leaves the area as direct surface runoff, and a little percolates to augment the dissected groundwater aquifers (Choubey et al. [1999\)](#page-15-14) that ultimately return to the surface in the form of springs.

#### **Geological setup and geothermal activity**

The Himalayan tectonic structure is a blend of complex processes that function on diferent time scales. It conglomerates the Himalayan orogen, active foreland basin and major depositional systems like the Indus and the Bengal fans. The northwest and northeast Himalayan syntaxes are the marked bands of strike on both ends of the Himalaya (Tiwari [2014\)](#page-17-3). The orogen is diferentiated by ITSZ in the north, the Chaman fault in the west, the Sagging fault in the east and the Main Frontal Thrust (MFT) in the south (LeFort [1975\)](#page-16-15). South-north division of the Himalaya starts from the Shivaliks; proceeding to the Lesser, Higher and Tethys Himalaya; and fnally the Indus Suture Zone, which connects with the Trans-Himalaya (Auden [1937\)](#page-15-15). These geographical and geological zones continue along the entire Himalayan arc (Gansser [1964](#page-15-16); LeFort [1975](#page-16-15)). Thus, Garhwal Himalaya provides a perfect site for learning the diverse geological processes related to mountain building. Many researchers in recent years have been motivated by the potential of the region to decipher the feedback procedures between lithospheric deformation and atmospheric circulation as a driver of global climate change (e.g. Molnar [1987;](#page-16-1) Harrison et al. [1992](#page-16-16), [1998;](#page-16-17) Royden et al. [1997](#page-16-18); Ramstein et al. [1997](#page-16-19); Tapponnier et al. [2001;](#page-17-8) Beaumont et al. [2000](#page-15-17); Yin et al. [2002](#page-17-9)). This study covers the geothermal systems of Garhwal Himalaya in Alaknanda, Bhagirathi and Yamuna valleys along the MCT zone (Heim and Gansser [1975](#page-16-20)).

The Higher Himalayan Crystalline (HHC) rocks along these river valleys, separate from the Tethyan Himalaya to the north by the South Tibetan Detachment (STD) and from the Lesser Himalaya to the south by the southern strand of the MCT (Munsiari Thrust, or MCT-I, Fig. [1\)](#page-2-0). The HHC contains high-grade gneisses, metabasite, quartzite, schist and granite, with limited carbonate and calc-silicate rocks



<span id="page-2-0"></span>**Fig. 1** Geological map of the Garhwal Himalaya with details of sampling locations of geothermal systems and river waters

(Aggarwal and Kumar [1973](#page-15-18)). The metamorphic grade of the HHC increases from biotite in the south with chlorite phyllite, chlorite schists and metabasic rock south of the MCT-I to kyanite and sillimanite bearing rocks north of the Vaikrita Thrust (or MCT-II; Sachan et al. [2001\)](#page-16-21). It is the response of an inclined fault that got detached from the upper crust along a sub-horizontal zone at an inclination of 30° to 40° northward. It has also sufered a thrusting up to a depth of ~ 125 km which brought basement rocks up to the surface through ductile deformation (Roy and Valdiya [1988](#page-16-22)). Nearby the MCT in the Lesser Himalayan zone, the metamorphic and associated granite rocks are found severely deformed and compressed (Bhattacharya [1985\)](#page-15-19). The majority of geothermal systems are around the MCT zone (fanked by MCT-I and MCT-II) in the Garhwal Himalaya, a region of high heat flow  $(130 \pm 30 \text{mW/m}^2)$  and high-temperature gradient ( $60^{\circ} \pm 20^{\circ}$ C/km) that contrasts with the low heat flow  $(41 \pm 10 \text{mW/m}^2)$  and the low-temperature gradient  $(17^{\circ} \pm 5^{\circ}$ C/km) of the lower Himalaya, south of the MBT (GSI [1991\)](#page-15-20). Hence, sub-surface temperatures are high in the MCT zone, possibly due to the presence of partial melts and their associated fuids or perhaps simply due to coupled erosion-rock uplift (Derry et al. [2009\)](#page-15-2). The electrical resistivity below 10Ω-m at 10–20-km depth in the MCT zone, along both the Bhagirathi and Alaknanda valleys, is thought to represent either partial melt or metamorphic fuids (Rawat et al. [2014](#page-16-23)) at moderate to high temperatures, thus providing

a continuous heat source for the geothermal systems in the region.

The geothermal systems of Garhwal Himalaya largely fall in the central crystalline zone consisting of the Munsiari formation of the lower Himalayan crystalline sequence (LHCS) followed by the Vaikrita group of the greater Himalayan sequence (GHS) including Joshimath, Pandukeshwar and Badrinath formations. The Munsiari formation includes garnet-bearing mica schists, calc-silicate lenses and quartzite (Gururajan and Choudhuri [1999](#page-16-24)). The mineral assemblage within the Munsiari formation includes quartz, biotite, plagioclase, garnet, staurolite, muscovite, graphite and andalusite. The MCT in the Garhwal region is the ductile shear zone between the micaceous quartzites and chlorite-biotite schists of the Munsiari formation and the kyanite gneisses of the Joshimath formation. The Joshimath formation has a sequence of pelitic gneisses with schistose interlayers. It has a mineral assemblage of quartz, biotite, plagioclase, garnet, staurolite, muscovite, kyanite, chlorite, calcite, graphite, titanite and opaques (rutile, ilmenite). The Joshimath formation grades from metapelites to meta-arkose of Pandukeshwar formation. The Pandukeshwar formation consists of meta-arkose and quartzite. The meta-arkose is primarily medium- to fne-grained and is interlayered with kyanite-bearing schist (Paul [1998](#page-16-25)). Mineral assemblages in this formation include quartz, K-feldspar, plagioclase, muscovite, biotite, garnet, chlorite and opaques (rutile,

ilmenite, Cr-rich spinel). The Badrinath formation primarily consists of migmatitic metapelites and calc-silicates that grade upward from the meta-arkose of the Pandukeshwar formation. The Badrinath formation is bounded at the top by the STD and structurally overlain by the unmetamorphosed Martoli formation of the Tethyan Sedimentary Series. The metapelites of the Badrinath formation have a similar mineral assemblage to the Joshimath formation with the addition of sillimanite (quartz, biotite, plagioclase, garnet, staurolite, muscovite, kyanite, sillimanite, cordierite, chlorite, calcite, graphite, rutile and ilmenite) (Spencer et al. [2012\)](#page-17-10). Calcsilicate layers generally contain calcite, diopside, quartz, hornblende, biotite, scapolite, actinolite and tremolite (Paul [1998;](#page-16-25) Spencer et al. [2012](#page-17-10)). Migmatitization of the Badrinath formation occurs  $\sim$  16 km up the structural section from the MCT-II and increases upward to the STD. Migmatite stringers coalesce in the upper Badrinath formation where large leucogranite bodies intrude the surrounding metasediments. The leucogranite bodies consist of quartz, K-feldspar, plagioclase, muscovite, biotite, garnet, tourmaline, sericite (secondary) and chlorite (secondary).

Following the pioneering works of Auden ([1937\)](#page-15-15) and Heim and Gansser ([1975](#page-16-20)), numerous geological studies have been carried out along the Yamuna, Bhagirathi, and Alaknanda valleys of the Garhwal Himalaya (Valdiya, [\(1980;](#page-17-11) [1981;](#page-17-12) [1988\)](#page-17-13); Thakur and Rawat ([1992\)](#page-17-14)). Details of occurrences, discharge type and lithological formation of geothermal systems are given in Table S1.

#### **Methodology**

A total of twenty (20) samples of geothermal systems and ten (10) samples of river water associated with these geothermal systems were collected during 2010, 2012, 2013 and 2017 for the isotope (stable and radiogenic) and geochemical characteristics of the geothermal systems (Fig. [2A](#page-4-0)–[F](#page-4-0)). Besides, glacier ice (01), snow (03), high-altitude rainfall (04) and meltwater (04) samples were also collected from Dokriani Glacier during 2010, 2013 and 2014. The snowpit profle was conducted in May 2013 in the ablation zone of Dokriani Glacier (Fig. [2G](#page-4-0)). Geothermal spring water samples were collected close to their mouth, while river/ glacial meltwaters were collected from the middle of the stream (Fig. [2H](#page-4-0)). The altitudinal range for all samples varies from 800 to 4068 m above sea level (asl). All water samples were fltered and stored in pre-cleaned HDPE amber colour bottles (Tarson®). One aliquot of the fltered water samples was poisoned by adding a 10 μL saturated solution of  $HgCl<sub>2</sub>$  (Sigma-Aldrich®) to prevent biological growth for stable isotope analyses. The physical parameters of all water samples were measured in situ (Fig. [2\)](#page-4-0), including the temperature ( $\rm{^{\circ}C}$ ), pH and electrical conductivity ( $\mu$ S/cm, EC) as per established methods (Dalai et al. [2002](#page-15-21); Tiwari

et al. [2016a, b](#page-17-5)) using the multi-electrode probes (The LaMotte pH 6 Series, Con6 meter) during the sampling with an average precision of  $\pm$  0.01 for conductivity,  $\pm$  0.05 for pH and  $\pm 0.05$  °C for temperature.

# **Chloride (Cl−) and dissolved silica (SiO2) analysis**

Chloride (Cl−) of the samples was analysed using ion chromatography (Dionex series ICS-5000) at the Wadia Institute of Himalayan Geology (WIHG) Dehradun. The calibration of the instrument and analysis of samples was done following the procedure of Tiwari et al. ([2016a](#page-17-5), [b](#page-17-2), [2020](#page-17-1)). The precision based on repeat analysis was  $\pm$  5% (1SD). Further, the dissolved silica was analysed in an un-acidifed, fltered aliquot of the samples using a quadrupole inductively coupled plasma-mass spectrometer (ICP-MS) at WIHG (Tiwari [2014](#page-17-3)). The reproducibility of the measurement for dissolved silica was  $\pm$  5% (1SD).

# **Isotopic analysis (δ18O ‰vsmow,δD ‰vsmow and 87Sr/86Sr ratio)**

The measurement of the stable oxygen isotope  $(\delta^{18}O)$  was carried out at the Isotope Ratio Mass Spectrometry Laboratory (IRMS) at WIHG following the established method by Tiwari et al. ([2016a](#page-17-5), [b](#page-17-2), [2018,](#page-17-15) [2020](#page-17-1)), Verma et al. [\(2018\)](#page-17-16) and Kumar et al. ([2018](#page-16-26)). The analytical procedure for the measurements of  $\delta D$  is almost similar to that of  $\delta^{18}O$  except for the equilibration method where a platinum stick is placed into the vial before fush flling (Tiwari et al. [2016a](#page-17-5), [b](#page-17-2); Verma et al. [2018;](#page-17-16) and Kumar et al. [2018\)](#page-16-26). IRMS was calibrated with primary standards of GISP and VSMOW supplied by IAEA for measurements at regular intervals (Table S2, Craig [1961b](#page-15-22)). The reproducibility of measurement was 0.06‰ and  $1\%$  (1SD) for  $\delta^{18}$ O and  $\delta$ D, respectively.

For Sr isotopic measurement of water samples, Sr specifc resin (Eichrom) was used. The fltered water sample was loaded on the Sr specifc resin in a clean chemistry lab. Milli-Q water was used to elute the pure Sr fraction. This fraction was dried and then made up using  $3 N HNO<sub>3</sub>$ (Chatterjee and Singh [2012\)](#page-15-23). Sr isotopic measurement of the samples was analysed using Multiple Collector ICP-MS (Neptune Plus, Thermo Fisher Scientifc) in static mode. Instrumentation mass bias correction was done using the  $88$ Sr/ $86$ Sr ratio (typical value: 8.375209 with exponential law). The isobaric interferences of Rb and Kr are corrected by measuring 85Rb and 83Kr using natural abundances. However, the signal of  ${}^{85}Rb$  and  ${}^{83}Kr$  was weak. A total of 40 cycles for each measurement were run with an integration time of about 8.3 s per cycle. The total measurement time was approximately 6 min for each run.

Details of all sample results are given in Table [1,](#page-5-0) whereas that of glacier melt/snow samples collected in diferent years and cold springs are shown in Tables [2](#page-6-0) and S3. The pH of geothermal systems varied from 5.9 to 8.6 and is similar to the surface waters like glacier meltwater/fresh snow (6 to 8.6), natural springs (6.8 to 7.6) and headwaters of Ganga (7.3 to 8.2). Therefore, it indicates the identical source of water in the region (Fig. [3A](#page-7-0)). The mouth temperature of the water in geothermal systems varies from 26 to 93 °C. The temperature range for surface waters (glacier meltwater, cold springs and river waters) was 2 to 21 °C (Tables [1](#page-5-0) and [2](#page-6-0) and S3 and Fig. [3B\)](#page-7-0). Electrical conductivity (EC) of waters in geothermal systems varies from 340 to 4950 μS, while that for river waters, snow/glacier meltwater and cold springs varies from 59 to 271

**Results and discussions**

<span id="page-4-0"></span>**Fig. 2** Field photographs showing (**A**) Alaknanda at Badrinath, (**B**) release of the jet stream at Tapovan geothermal spring, (**C**) sampling at Vishnuprayag, (**D**) feld measurements of physical parameters at Helang, (**E**) vent of geothermal system at Gangnani, (**F**) feld measurements at Suryakund, Yamnotri, (**G**) snow pit sampling, and (**H**) temporary gauging site at Dokriani Glacier





<span id="page-5-0"></span>**Table 1** Details of samples collected and their in situ and laboratory measurements



<span id="page-6-0"></span> $\underline{\textcircled{\tiny 2}}$  Springer



<span id="page-7-0"></span>**Fig. 3** Distribution and variability of a pH, b temperature, c electrical conductivity and d chloride in diferent waters of Garhwal Himalaya

μS, 5 to 122 μS and 24 to 311 μS, respectively (Fig. [3C\)](#page-7-0). The EC of geothermal systems is higher by an order of magnitude as they are expressions of rock-water interaction at moderate to high temperatures.

#### **Variability in chloride (Cl−) and dissolved silica (SiO2)**

Hydro-geochemical analysis of geothermal systems is extensively applied as a fundamental method in distinguishing geothermal reservoirs and their evolution (Fournier [1981;](#page-15-24) Nicholson [1993\)](#page-16-27). Concentrations of chloride and dissolved silica of geothermal systems, river waters, cold springs and glacial meltwater are available in Tables [1](#page-5-0) and [2](#page-6-0) and S3. The concentration of Cl − in geothermal systems, river waters and cold springs varies from 95 to 21,986 μE/L, 7 to 40 μE/L and 6 to 274 μE/L, respectively, and the average is 3274 μE/L, 20 μE/L and 105 μE/L, respectively. The concentration of Cl− in fresh snow and glacial meltwater varies from 2 to 38 μE/L, with an average of 10 μE/L (Fig. [3D](#page-7-0)). Cl− is a major soluble ion of natural waters and does not precipitate except under evaporation and freezing processes. These conductive ions are derived from dissolved salts and inorganic materials such as alkalis, chlorides, sulphides and carbonate compounds (Miller et al. [1988](#page-16-28)). The high concentration of Cl − indicates rock-water interaction in the deep reservoir, where conductive cooling and mixing are low. Low Cl− values may specify the boundary and modern underground water interference to the geothermal system (Nicholson [1993](#page-16-27)). The observed variation in  $\delta^{18}$ O of all geothermal systems is related to their altitude, while the concentration of Cl− is relatively the same for all these systems. The variations in the  $\delta^{18}O$ and chloride (Cl−) are not by the efect of diverse evapo-concentrations or steam separation and are from the elevation difference of geothermal water recharge and sources of dissolved Cl−, respectively (Tan et al. [2014](#page-17-4); Tiwari et al. [2016a](#page-17-5)).

Dissolved silica  $(SiO<sub>2</sub>)$  is a vital component of all geothermal fuids. The solubility of silica polymorphs (amorphous silica, opal, quartz, cristobalite and chalcedony) and other silicate minerals controls its concentration. The abundance of dissolved  $SiO<sub>2</sub>$  in geothermal systems and river waters varies from 380 to 2040 μE/L and 159 to 255 μE/L, respectively. The average dissolved  $SiO<sub>2</sub>$  in geothermal systems and river waters is 1399 µE/L and 205 µE/L, respectively (Table [1](#page-5-0)). The results strongly advocate that the surface waters infltrate deep into the ground, possibly interacting with the silicates in the clayey level or minerals of volcanic rocks (Fig. [4A](#page-9-0)).

# **Variability in stable isotopic data (δ18O‰, δD ‰) and identifcation of the water source**

Stable isotopes of water have been used as proxies to unearth the recharge source of geothermal systems in this study. The ratios of stable isotopes of  $(\delta^{18}O, \delta D)$  of all the water samples are shown in Tables [1](#page-5-0), [2](#page-6-0) and S3, whereas their variations are given in Fig. [4B and C.](#page-9-0) The ratio of stable isotopes of water ( $\delta^{18}O$ ,  $\delta D$ ) has an average of  $-9.4\%$  and  $-65.2\%$  in geothermal systems,  $-12.4\%$ and –80.2‰ in river waters and –12.6‰ and –88.7‰ in the rain, fresh snow, glacial ice and meltwater of Dokriani Glacier. The cold springs of the Garhwal region have an average of –7.4‰ and –53.1‰. Similar values of stable isotopes of water in high-altitude rainfall, snowfall and glacial meltwater have been reported for Chorabari and Dokriani glaciers (Kumar et al. [2018](#page-16-26); Verma et al. [2018](#page-17-16)). The range of stable isotopes of water in geothermal systems is covered by diferent waters like high-altitude rainfall, snowfall and glacial meltwater.

Geothermal systems may integrate waters from diverse sources, which could be resolved using stable isotopes of water. Friedman ([1970](#page-15-25)) first reported the dominance of the meteoric component in geothermal systems by observing δD values; however, a better understanding came through the combined measurements of  $\delta^{18}$ O and  $\delta$ D (Craig [1963](#page-15-26)). The equilibrium fractionations describe the relationship between  $\delta^{18}$ O and  $\delta$ D composition in natural waters (Craig [1961a](#page-15-27); Dansgaard [1964](#page-15-28); Rozanski et al. [1993](#page-16-29)). Several studies led to the establishment of a Global Meteoric Water Line (GMWL) that is a relation between  $\delta^{18}O$  and δD of precipitation from various parts of the world, having a slope of 8 and intercept of 10, which is described as " $\delta D = 8 * \delta^{18}O + 10$ ". In contrast,  $\delta^{18}O$  and  $\delta D$  relationship at a regional or local scale is known as the Local Meteoric Water Line (LMWL). The intercept and slope may vary as a function of the location, seasonality and occurrences of extreme events. The "deuterium excess" or "d-excess" in precipitation is the intercept of the GMWL. It provides a measure of non-equilibrium efects (Dansgaard [1964\)](#page-15-28) and indicates the environmental circumstances during phase transfer under a non-equilibrium environment.

The δD values of geothermal systems lie close to the local meteoric water, while  $\delta^{18}$ O values are enriched. This shift in the  $\delta^{18}$ O but not in  $\delta$ D of meteoric water can help understand the hydrothermal interactions at depths. Geothermal systems and volcanic landscapes have fractures and crevices through which rainfall penetrates the earth and goes to a certain depth prior to interacting with hot rocks and getting back to the surface. During this process, oxygen from the meteoric water endures exchange reactions with source rocks (crystalline igneous/carbonates), having enriched  $\delta^{18}O$  in contrast with meteoric waters. Therefore, it would augment the enrichment of  $\delta^{18}O$  in the water and depletes the  $\delta^{18}O$  in the rock. However, there is very little hydrogen in igneous and carbonate rocks for exchange during this process, and the δD composition of water is unaltered. In such a situation, it forces the water line to the right side of the LMWL (Witcher et al. [2004](#page-17-17)).



<span id="page-9-0"></span>**Fig. 4** Distribution and variability of a dissolved silica (SiO<sub>2</sub>), bδ<sup>18</sup>O, c δH and d<sup>87</sup>Sr/<sup>86</sup>Sr ratio in different waters of Garhwal Himalaya

Therefore, the  $\delta^{18}O$  and  $\delta D$  relationship of geothermal waters is useful to study the altitude effect and source of water in geothermal systems.

The LMWL for geothermal systems, river water, meltwater, rainfall and fresh snow in the study region lies over or close to the GMWL with varied slope and intercepts (Fig. [5](#page-10-0)). D-excess in geothermal systems in the study area ranges from 6.9 to [1](#page-5-0)9.4, with a mean of  $13.2 \pm 6.2$  (Tables 1 and S4). The d-excess of these systems is quite similar to glacial meltwater, high-altitude rainfall and snowfall in UGB (Ramesh and Sarin [1992](#page-16-30); Bartarya et al. [1995](#page-15-29); Dalai et al. [2002;](#page-15-21) Kumar et al. [2018;](#page-16-26) Verma et al. [2018](#page-17-16)). The slope and intercept for geothermal systems are similar to glacial meltwater, highaltitude rainfall and snowfall in UGB. The slope, intercept and d-excess of cold springs, Yamuna and Indus Rivers are different (Fig. [6\)](#page-11-0), because of the different climates, location and altitude of these systems. Therefore, the slope, d-excess and intercept of geothermal systems indicate that high-altitude rainfall, snowfall and meltwater from glaciers are the foremost source of recharge of these geothermal systems. Although the geothermal fuids exhibit meteoric origin, chemical compositions and variations indicate the likelihood of a small (5–10%) but noteworthy addition of the magmatic or deeper component (Nicholson [1993](#page-16-27); Barbier [2002](#page-15-30); Tiwari [2014](#page-17-3)).

## **Identifcation of the source of the geothermal fuid using 87Sr/86 Sr ratio**

The Himalayan Rivers contribute signifcantly to the Sr isotope mass balance of the modern oceans and have high Sr and elevated  ${}^{87}Sr/{}^{86}Sr$  among the world rivers (Edmond [1992](#page-15-31)). The chief source of radiogenic Sr fux in Himalayan Rivers is from processes like weathering of diferent silicate phases and radiogenic carbonates (Palmer and Edmond [1989;](#page-16-31) Derry and France-Lanord [1996](#page-15-32); Quade et al. [1997](#page-16-32); Blum et al. [1998](#page-15-33); Singh et al. [1998;](#page-17-18) Galy and France-Lanord [1999](#page-15-34); Krishnaswami [1999](#page-16-33); English et al. [2000\)](#page-15-35). Radiogenic isotope  $(^{87}Sr/^{86}Sr)$  ratios of all the water samples are given in Table [1,](#page-5-0) and their variations are shown in Fig. [4D](#page-9-0). The radiogenic isotope ratio of strontium  $({}^{87}Sr/{}^{86}Sr)$  varies from 0.73 to 1.44‰, with an average of  $0.83\%$  in geothermal systems. In the associated river waters, the ratio varies from 0.71 to 0.73‰, with an average of 0.72‰ (Fig. [4D](#page-9-0)), indicating diferent water–rock reactions between them. These results reveal that the river waters did not infltrate deep into the thermal reservoirs and are not the recharge source of geothermal systems (Tan et al. [2014;](#page-17-4) Zhao et al. [2003](#page-17-19)). Several previous studies suggest that geothermal systems contribute a signifcant amount of radiogenic strontium to the Himalayan Rivers and deliver up to 30% of the Sr fux to rivers (Evans et al. [2001](#page-15-36)). Therefore, the Earth's internal fuids/magma/heat source interacts with the water at depths for transferring the heat and radiogenic  ${}^{87}Sr/{}^{86}Sr$  to the water in geothermal systems.

#### **Circulation of water in geothermal systems**

The recharge source of geothermal systems originates at a high altitude, which is beneficial to understand the movement of geothermal water and estimate the recharge altitude.

<span id="page-10-0"></span>**Fig. 5** Meteoric water line for diferent waters of Garhwal Himalaya and comparison with **GMWL** 



<span id="page-11-0"></span>



The isotopic composition of precipitation varies with the altitude of the sampling location (Yurtsever and Gat [1981](#page-17-20); Clark and Fritz [1997\)](#page-15-12) and is known as the "altitude efect". As air masses lift orographically, they cool and cause precipitation which is enriched in heavier isotopes. The next phase of precipitation condensing from the residual cloud mass at a higher altitude would deplete in heavy isotopes; this process progressively increases with altitude. Therefore, the precipitation history, topography and perceptible moisture remaining in the cloud control the altitude efect. For  $^{18}$ O, the depletion varies between –0.15 and –0.5‰ per 100 m rise in altitude, with a corresponding decrease of –1 to  $-4\%$  for  $^2H$  (Yu et al. [1984](#page-17-21); Schotterer et al. [1997](#page-16-34); Wright et al. [2001;](#page-17-22) Blasch and Bryson [2007\)](#page-15-37). Rayleigh distillation manifests continental and altitude efects on a large spatial scale due to cloud migration and the continuous withdrawal of water from the ascending clouds. Krishnamurthy and Bhattacharya ([1991\)](#page-16-35) have observed the continental effect in precipitation over northern India. The rivers also refect the altitude efect associated with precipitation, as precipitation is the dominant water source in rivers. However, river water is likely to be more depleted relative to precipitation at that site as it represents a mixture of various waters from streams draining from higher altitudes up to the site.

Geothermal systems of the Garhwal region indicate an altitude effect in  $\delta^{18}$ O and  $\delta$ D. The altitude of the geothermal systems of Garhwal Himalaya varies from ~ 1160 to 3089 m asl, and the value of  $\delta^{18}$ O varies from -7.5‰ to –13.1‰ (Table [1](#page-5-0)). The altitude efect for the geothermal systems and the UGB is  $-0.2 \pm 0.025\%$  per 100 m and  $-0.2 \pm 0.006\%$  per 100 m (Fig. [7](#page-12-0)). These lie within the values reported in earlier studies in the Garhwal Himalaya (Table [3](#page-12-1)). Some geothermal systems have an altitude higher than the River and have depleted isotopic ratios than the river waters (Table [1](#page-5-0)). The geothermal systems which have a lower elevation than the river show enriched ratios although rain and glacial meltwater from an altitude of 3810 m asl demonstrate similar isotopic values as water in geothermal systems. The results for the UGB, Gaula and Yamuna rivers in Garhwal Himalaya (Ramesh and Sarin [1992](#page-16-30); Bartarya et al. [1995;](#page-15-29) Dalai et al. [2002\)](#page-15-21) and Seti and the Kali Gandaki watersheds in Nepal (Garzione et al. [2000a](#page-15-38), [b\)](#page-15-39) also show the altitude effect in  $\delta^{18}$ O and  $\delta$ D values of rivers (Table [3](#page-12-1)). The precipitation collected from 11 sites between 250 and 3250 m asl at Mount Meager, a Quaternary volcano in western British Columbia, shows an altitude efect of –0.25‰ per 100 m (Clark et al. [1982](#page-15-40)) which provided evidence for the recharge environment of the geothermal groundwater. A gradient of  $-0.2\%$  per 100 m for  $\delta^{18}$ O prevails in the Jura Mountains of northern Switzerland (Siegenthaler and Matter [1983\)](#page-16-36). A schematic physical model of the recharge process for geothermal systems of the Garhwal region is shown in Fig. [8](#page-13-0). The top panel of the physical model (Fig. [8A\)](#page-13-0) represents the process of infltration of the meteoric waters (rainfall, snow and glacier melt) from high altitude into the ground through the faults/thrusts of the region and subsequent heating from the underlying heat source in the MCT zone and is released as geothermal springs. The Moho is the boundary between the crust and the mantle in the earth. This is a depth where seismic waves change velocity and there is also a change in chemical composition. The resistivity data (Fig. [8B\)](#page-13-0) in the lower panel of the physical model indicates very low resistivity values in the MCT zone at comparatively low depths (5–20 km) indicating the presence of partial melt

<span id="page-12-0"></span>



<span id="page-12-1"></span>**Table 3** Altitude efect in geothermal systems, cold springs, rainfall and rivers in the Himalayan region



or metamorphic fuids (Caldwell et al. [2013\)](#page-15-41). These fuids are the primary heat source for the geothermal systems and provide the thermal gradient for heating the meteoric waters. Therefore, the meteoric waters infltrate to a shallow depth and are heated and released as geothermal waters.

# **Geothermal systems: a permanent source of Earth's internal heat**

The data of the mouth temperature of the geothermal systems of Garhwal Himalaya since 1975–1994 (Geological Survey of India) and 2010–2016 (this study) signify that the mouth temperatures of these systems remain the same over the last four decades with an error of  $< 5\%$  (Fig. [9](#page-13-1) and Table S5). The extent of dissolution and precipitation of diferent substances and their solute characteristics depend upon the temperature and water–rock interactions with thermal fuids extracted from the deep reservoir. Hence, it is essential to understand the temperature-induced process and infer the character of the thermal reservoirs. Field observations, chemical composition and physical parameters of these geothermal systems show no temporal variation in <span id="page-13-0"></span>**Fig. 8** A physical model depicting the process of recharge of geothermal systems (**a**) and resistivity data (**b**) for Garhwal Himalaya



<span id="page-13-1"></span>**Fig. 9** Variation in the water temperature at the mouth of diferent geothermal systems of Garhwal Himalaya

their constituent (Gupta and Agarwal [2001;](#page-15-8) Tiwari et al. [2016a\)](#page-17-5). Previous studies in other parts of the Himalaya also reveal similar results (Piao et al. [2010](#page-16-39); Meng et al. [2012](#page-16-40); Tan et al. [2014\)](#page-17-4), thus indicating that geothermal systems act as a profuse, continuous source of energy which is dependent on post-tertiary granite intrusions for the high thermal gradient (>100 °C/Km) and heat flow (>468 mW/m<sup>2</sup>) (Chandrasekharam and Chandrasekhar [2000](#page-15-42)).

Further, the discharge of geothermal systems has remained more or less unchanged over the period between

<span id="page-14-0"></span>



1975 and 2016 (Fig. [10\)](#page-14-0), with small variations observed due to changes in the season of sampling or increased snow and glacier meltwater in the Garhwal region. Therefore, the recharge of geothermal systems is a continuous process, and availability of meteoric waters like high-altitude precipitation (rainfall and snowfall) and glacier melt will keep these geothermal systems healthy. It is a comparatively clean and renewable energy resource and hence becoming a preferred choice for alternative energy. In India, the estimated potential to generate  $\sim 10,600$  MWe of power (Craig et al. [2013\)](#page-15-43) using the latest technologies in geothermal systems is still unutilised.

## **Conclusions**

This study concludes that the diferent geothermal systems of Garhwal Himalaya have high electrical conductivity (EC) by an order of magnitude than the background due to rockwater interactions at moderate to high temperatures. The enriched Cl<sup>−</sup> in the geothermal systems may be derived from the chloride phase present in the metamorphic crustal fuid. The radiogenic  ${}^{87}Sr/{}^{86}Sr$  data reveals that river water is not the recharge source of geothermal waters.

Based on the isotopic composition ( $\delta^{18}$ O and  $\delta$ D), slope, d-excess and intercept of the LMWL of geothermal waters, it is evident that meteoric waters (rainfall, snow and glacier melt) are the principal recharge source of geothermal systems in Garhwal Himalaya. Therefore, it is important to

monitor the seasonal and annual precipitation (rainfall and snowfall) and the state of Himalayan glaciers to understand the sustainability of these geothermal systems for utilisation of the communities living in downstream.

The discharge and mouth temperature of the water of these geothermal systems remains unchanged, thus indicating that these springs are a continuous source of energy and their potential can be harnessed for the generation of electricity and internal heating of buildings in the vicinity as done by several Scandinavian countries. Further, various studies have indicated the presence of heavy metals and carcinogens in the waters of the geothermal systems; thus, these waters should be carefully utilised for domestic and agricultural purposes.

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## **Declarations**

**Conflict of interest** The authors declare that they have no competing interests.

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