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# Geochemical Data from Groundwater at the Proposed Dewey Burdock Uranium In-Situ Recovery Mine, Edgemont, South Dakota

By Raymond H. Johnson



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U.S. Department of the Interior  
U.S. Geological Survey

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# Geochemical Data from Groundwater at the Proposed Dewey Burdock Uranium In-Situ Recovery Mine, Edgemont, South Dakota

By Raymond H. Johnson

## Abstract

This report releases groundwater geochemistry data from samples that were collected in June 2011 at the Dewey Burdock proposed uranium in-situ recovery site near Edgemont, South Dakota. The sampling and analytical methods are summarized, and all of the data, including quality assurance/quality control information are provided in data tables.

## Introduction

Powertech Uranium Corporation (Powertech) has proposed to mine uranium at the Dewey Burdock site using in-situ recovery methods. The Dewey Burdock site is located in the southwestern region of the Black Hills of South Dakota (fig. 1). The uranium recovery license application by Powertech to the United States Nuclear Regulatory Commission (NRC) is publicly available and contains background information about the site along with technical details and baseline sampling data. The NRC application Web site is: <http://www.nrc.gov/materials/uranium-recovery/license-apps/dewey-burdock.html> (with detailed application documents under the “application documents” link). A brief summary of the site history is provided by Powertech at <http://www.powertechuranium.com/s/DeweyBurdock.asp>.

At the Dewey Burdock site, uranium occurs as roll-front ore bodies in several sandstone units of the Inyan Kara Group of Early Cretaceous Age. The Late Jurassic Morrison Formation underlies the Inyan Kara Group. In the vicinity of the mine site, the Inyan Kara Group is comprised of the Fall River Formation (upper unit) and the Lakota Formation (lower unit). The Lakota Formation is comprised of the Chilson and the Fuson Members, with the Fuson Member occurring between the Fall River and Lakota Formations. Uranium roll-front deposits occur in the Fall River Formation and the Chilson Member of the Lakota Formation. Other geologic units of interest for the study area are the surface alluvial aquifers and the Unkpapa aquifer underlying the Morrison Formation. The well location symbols on the maps in figures 2 through 4 are color coded to indicate the aquifer screened interval for each well.

The U.S. Geological Survey (USGS) collected 28 groundwater samples from monitoring wells (figs. 2 through 4) in and around the Dewey Burdock site during the last two weeks of June 2011. This sampling was completed with funding from the U.S. Environmental Protection Agency (USEPA) Region 8’s Regional Applied Research Effort (RARE) Program. USEPA is responsible for evaluating the site through its underground injection control program

(<http://www.epa.gov/region8/water/uic>) and Powertech has submitted a separate permit application to USEPA (see documentation in a link within the above USEPA Web site). While these new samples provide data on major ion and metal concentrations that overlap with Powertech's baseline sampling, the main intent of this sampling is to get isotopic measurements that will be used to better understand the hydrogeologic system.

## Sampling Methods

Groundwater samples were collected using either a peristaltic pump (used in shallow wells), a bailer (used when access problems were encountered), installed pump, or using a low-flow, submersible, stainless steel pump. Many of the wells are under artesian pressure and are flowing, with a valve assembly attached at the surface to control the flow (fig. 5). This same valve assembly was present in all wells with installed pumps. For wells with installed pumps or that were flowing, approximately three well bore volumes were purged prior to sampling.

A peristaltic pump was used in shallow 2-inch diameter wells where the water table was less than 27 feet below the top of the well casing. The sample tubing was placed approximately 1 foot above the bottom of the well and groundwater was purged until field parameters stabilized, which usually occurred within a few minutes. New polyethylene sample tubing was used for each well.

The submersible pump was a Geotech SS Geosub model by Geotech Environmental. The stainless pump was placed at approximately 50 feet below the water table and a drop tube assembling was lowered ahead of the pump to the bottom of the well ([http://www.geotechenv.com/pdf/ground\\_water\\_sampling\\_equipment/ss\\_geosub\\_wcontroller.pdf](http://www.geotechenv.com/pdf/ground_water_sampling_equipment/ss_geosub_wcontroller.pdf)). The tubing and pump were then pulled back so the intake was one foot from the bottom of the well. This allowed for direct sampling within the well screen and within the active groundwater flow zone using micropurge sampling. Groundwater was purged until field parameters stabilized, which usually occurred within a few minutes and was generally a function of meter stabilization (partially because of added pumping pressure) and not a change in groundwater conditions. New polyethylene sample tubing was used for each well.

A bailer was used in well 680 when the installed pump in the well failed after having purged three casing volumes. The well head was opened and the bailer was used to get water within the casing. A bailer was also used for well NBA, a newly installed well with a 2-inch diameter casing where the water table was below the limit of the peristaltic pump. A bailer was used to purge the well of three casing volumes before sampling. Sampling was completed using the peristaltic pump with new sample tubing to get water out of the bailer and through an inline filter.

For all samples, groundwater sampling was completed through a "T-valve" apparatus that split the flow into a flow through cell for field parameter measurement and a separate line for filling sample bottles (fig. 6). The sample line included an inline 0.45 micron filter that was used for all bottles except for tritium (sample NBA was filtered for tritium also, as the sediment load was high). Field parameter measurements of pH, specific conductance, temperature, dissolved oxygen, and oxidation/reduction potential were all done using a YSI 556 multiparameter meter that screwed directly into the flow through cell (see cover photograph), thereby eliminating any contact with the atmosphere.

Nine aliquots of water provided samples for (1) cations/metals, (2)  $^{234}\text{U}/^{238}\text{U}$  activity ratios, (3) anions, (4) dissolved organic carbon, (5) iron pairs ( $\text{Fe}^{3+}/\text{Fe}^{2+}$ ), (6) tritium, (7) stable isotopes ( $^{18}\text{O}$  and deuterium), (8) sulfur isotopes ( $^{34}\text{S}$ ), and (9) carbon isotopes ( $^{14}\text{C}$ ). Details on bottle type,

bottle size, rinsing, filtration, and preservative methods are listed in table 1. Any acids used for preservatives were ultra pure and made specifically for sample preservation. For bottles that were rinsed, rinsing was completed three times prior to filling the bottle.

**Table 1.** Bottle type and size, rinsing, filtration, and preservation for analytes.

[HDPE, high density polyethylene; mL, milliliter; L, liter;  $\mu\text{m}$ , micrometer;  $\text{HNO}_3$ , nitric acid;  $\text{HPO}_4$ , phosphoric acid;  $\text{HCl}$ , hydrochloric acid]

Analytes	Bottle Type	Bottle Size	Rinsing	Filtration	Preservative
Cations and dissolved metals	HDPE	30 mL	New bottle, rinse with sample water	0.45- $\mu\text{m}$	5 drops $\text{HNO}_3$ or to pH less than 2
$^{234}\text{U}/^{238}\text{U}$	HDPE	30 mL	New bottles, rinse with sample water	0.45- $\mu\text{m}$	5 drops $\text{HNO}_3$ or to pH less than 2
Anions	HDPE	30 mL	New bottle, rinse with sample water	0.45- $\mu\text{m}$	Keep cool
Dissolved organic carbon	Amber glass	125 mL	New bottles, cleaned and burned, do not rinse with sample water	0.45- $\mu\text{m}$	5 drops $\text{HPO}_4$ or to pH less than 2, keep cool
Dissolved iron species	Amber polyethylene	60 mL	New bottle, rinse with sample water	0.45- $\mu\text{m}$	5 drops $\text{HCl}$ or to pH less than 2, keep cool
Tritium	HDPE	500 mL	New bottles, rinse with sample water	None	None
Water Isotopes: $^{18}\text{O}$ and Deuterium	Borosilicate glass	60 mL	New bottles, rinse with sample water	0.45- $\mu\text{m}$	None
$^{34}\text{S}$	HDPE	125 mL	New bottles, rinse with sample water	0.45- $\mu\text{m}$	2 drops of $\text{HNO}_3$ to reduce biotic activity
$^{14}\text{C}$	Amber glass	1 L	New bottles, cleaned and burned, do not rinse with sample water	0.45- $\mu\text{m}$	Keep cool

## Analytical Methods

This section describes the analytical procedures completed on each of the nine water sample aliquots. The cations/metals sample was analyzed at the USEPA (Ada, Oklahoma) and at the USGS laboratories (Denver, Colorado). USEPA analyses were made using inductively coupled plasma – optical emission spectrometry (ICP-OES, Perkin-Elmer Optima 3300DV) using EPA Method 200.7 and inductively coupled plasma – mass spectrometry (ICP-MS, PQExcell, Thermo Elemental) using USEPA Method 6020. USGS analyses were made using ICP-MS at the USGS Mineral Resources Laboratory (Denver, Colorado) following the method described in Lamothe and others (2002).

The  $^{234}\text{U}/^{238}\text{U}$  activity ratios were analyzed by Michael Ketterer at the Northern Arizona University in Flagstaff, Arizona using a sector field Thermo X Series II quadrupole ICP-MS unit. Details on the analytical method can be found in file Appendix F.

Dissolved anions were analyzed by the USEPA laboratories (Ada, Oklahoma) using capillary electrophoresis with ultraviolet (UV) detection (USEPA Method 6500). Capillary ion electrophoresis is a free-zone electrophoretic technique optimized for the analysis of anions with molecular weights of less than 200 grams/mole. The anions migrate and are separated according to their mobility in the electrolyte when an electrical field is applied through the open tubular fused silica capillary.

Concentrations of dissolved inorganic carbon were measured by the USEPA laboratories (Ada, Oklahoma) with a Dohrmann DC-80 Carbon Analyzer (USEPA Method 9060A). Iron pairs ( $\text{Fe}^{3+}/\text{Fe}^{2+}$ ) were completed by David Fey at the USGS Mineral Resources Laboratory (Denver, Colorado) using the ferrozine method for iron species discussed in Bangthanh To and others (1999). Tritium analyses were completed by Robert Michel at the USGS Isotope Laboratory in Menlo Park, California using liquid scintillation counting with a detection limit of approximately 0.6 tritium units (TU) (similar to USEPA method 906.0). Oxygen- and hydrogen-isotopic ratios of water were analyzed using a high temperature conversion elemental analyzer linked to a continuous flow isotope ratio mass spectrometer (Finnigan Delta plus XP). These analyses followed the methods presented by Lu (2009). Sulfur isotopes ( $^{34}\text{S}$ ) were analyzed by Christopher Eastoe at the University of Arizona following the method of Coleman and Moore (1978). Additional analytical procedures for that laboratory can be found at [http://www.geo.arizona.edu/research/iso\\_analytical.html](http://www.geo.arizona.edu/research/iso_analytical.html).

## Data

All of the resulting data are provided in table 2. Sample identification numbers in table 2 match the well identification numbers used in Powertech permit application documents. Original labeling included “B” and “D” in the sample name for ease of sample identification (found in appendix files) by area, “B” for Burdock and “D” for Dewey, but were not included in the final data table (table 2).

The samples in table 2 have been organized by categories to reflect geologic units that the wells are screened in along with general area locations. These are purely initial categories used by the author for later use in interpretations.

## Quality Assurance/Quality Control

For all USEPA solution measurements, quality assurance tests involved duplicate samples, blanks, sample matrix spikes, calibration check standards, and second-source quality control samples. Data for the laboratory quality assurance/quality control (QA/QC) checks are included in the original data that can be found in the appendixes. Uranium isotope data also included internal laboratory checks that can be found in Appendix F. All other laboratories used typical internal reference standards for the appropriate analyses and QA/QC information is available upon request. No internal laboratory QA/QC issues were found.

In addition to the internal laboratory checks, three duplicate samples and five blanks were submitted as part of the QA/QC process. Data from these duplicates and blanks are reported in table 3. No QA/QC issues were found in any of the duplicate samples. Blanks for all of the different sampling conditions are represented (Geosub pump, peristaltic pump, and bailer

sampling). For metals, cations, and anions, a few blanks did show values above the detection limits, but generally these values were well below any sample results. Only one blank had an iron concentration that was high enough compared to the groundwater samples to be of concern. Iron in the bailer blank (B-VS4) was 3.72 µg/L, which is likely because of incomplete cleaning of the bailer. Cleaning the bailer was difficult because of limited access inside the bailer in addition to sample NBA (well sampled prior to blank) having a high silt content. As a result, any measured iron values below 4 µg/L, may not be accurate, especially for samples using a bailer. Sample data were not blank corrected.

## Acknowledgments

Funding from the US EPA made this data collection possible. Special thanks go to Richard Wilkin (USEPA, Ada, Oklahoma) for managing the USEPA contract and organizing the sample submittals to the appropriate USEPA laboratories. We are grateful for well access provided by individual land owners and through arrangements made by Powertech. Powertech employees, Wyatt Van Eaton and Mike Beshore provided invaluable assistance in accessing the wells and assisting with equipment logistics. Facilities for equipment storage and a convenient staging area were provided by Mark Hollenbeck. In addition, the groundwater sampling included assistance from Brian Zimmerman, Andrew Mahan, Valois Shea, Tanya Gallegos, and John Horton.

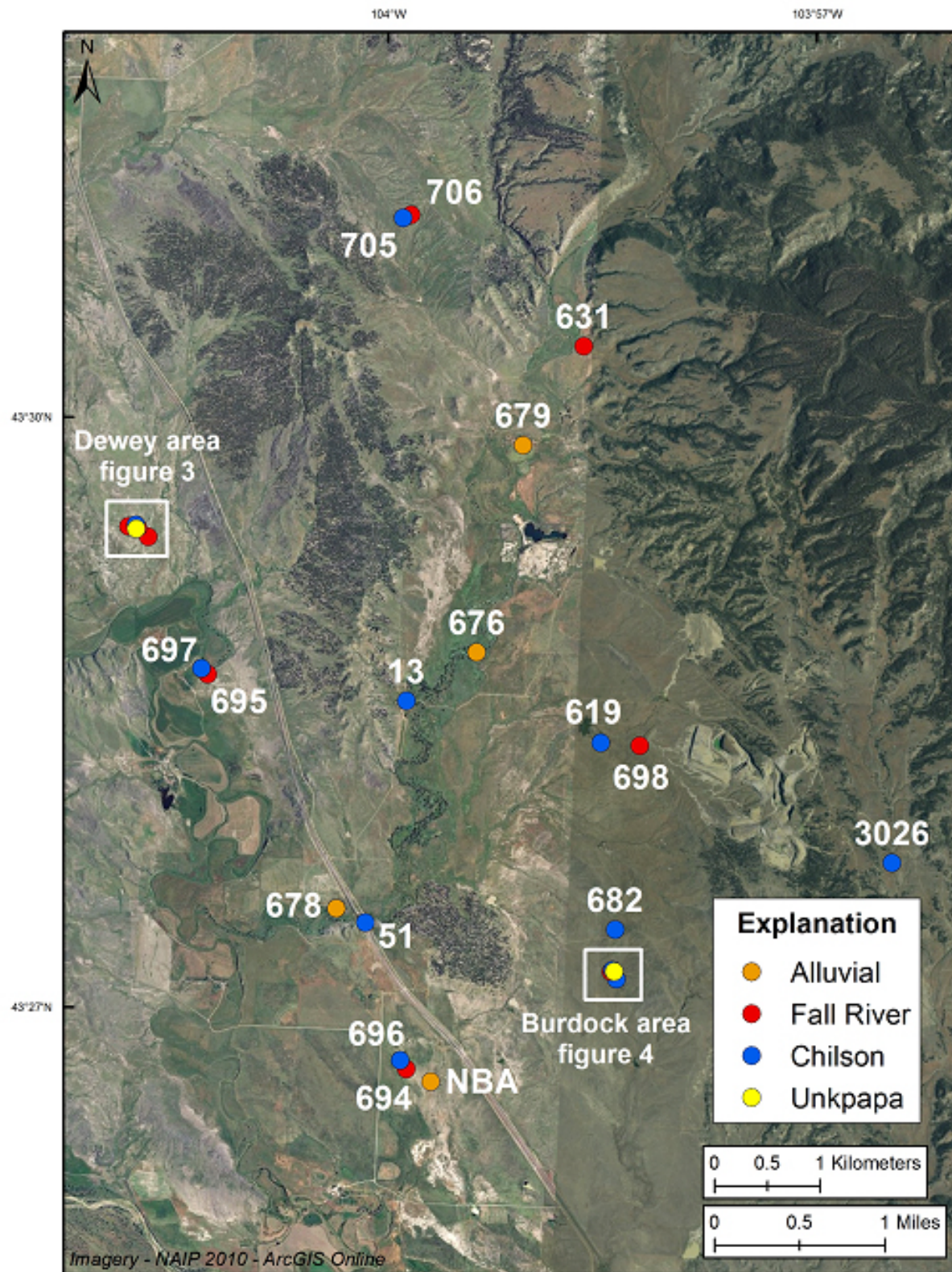
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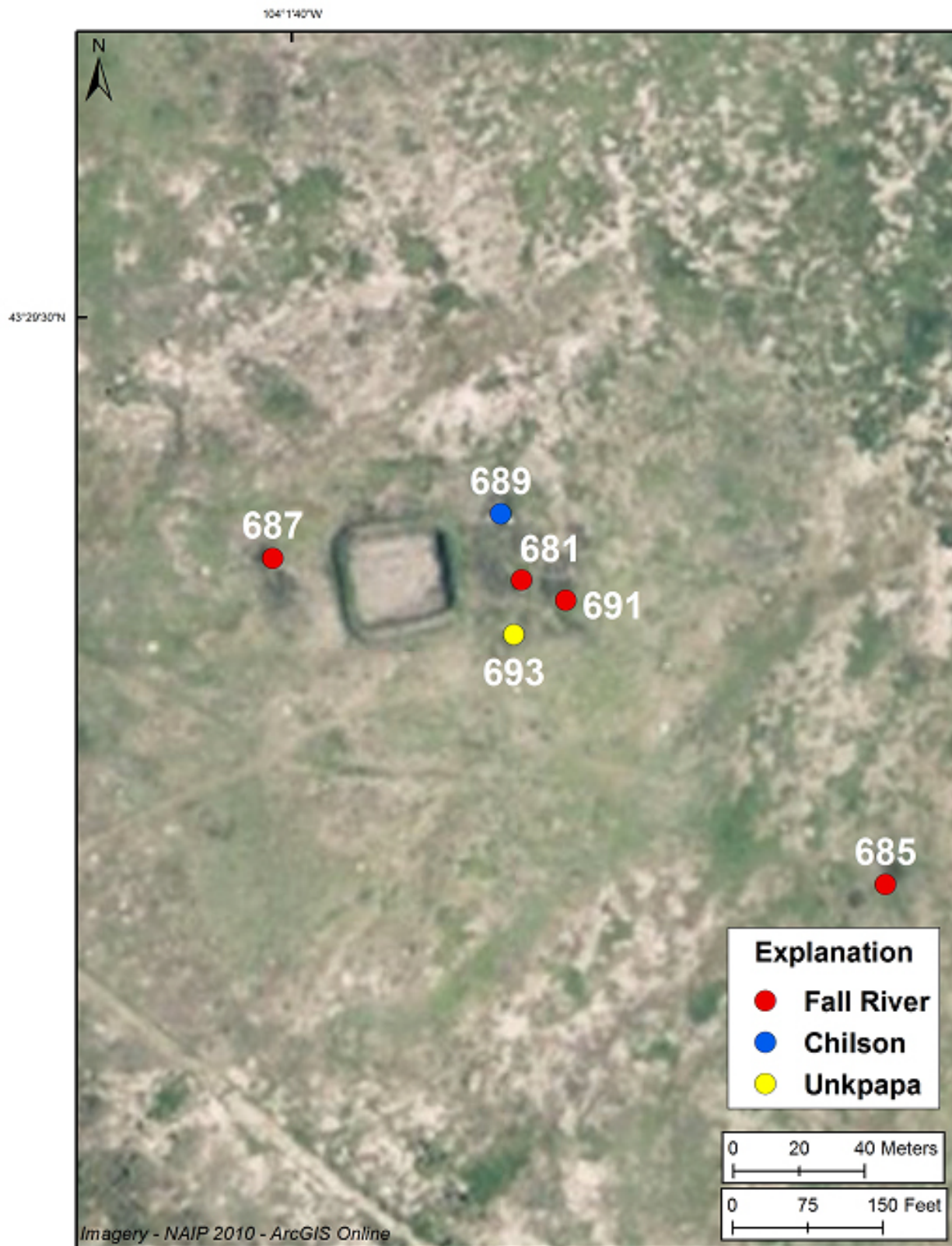


**Figure 1.** Location of study area.

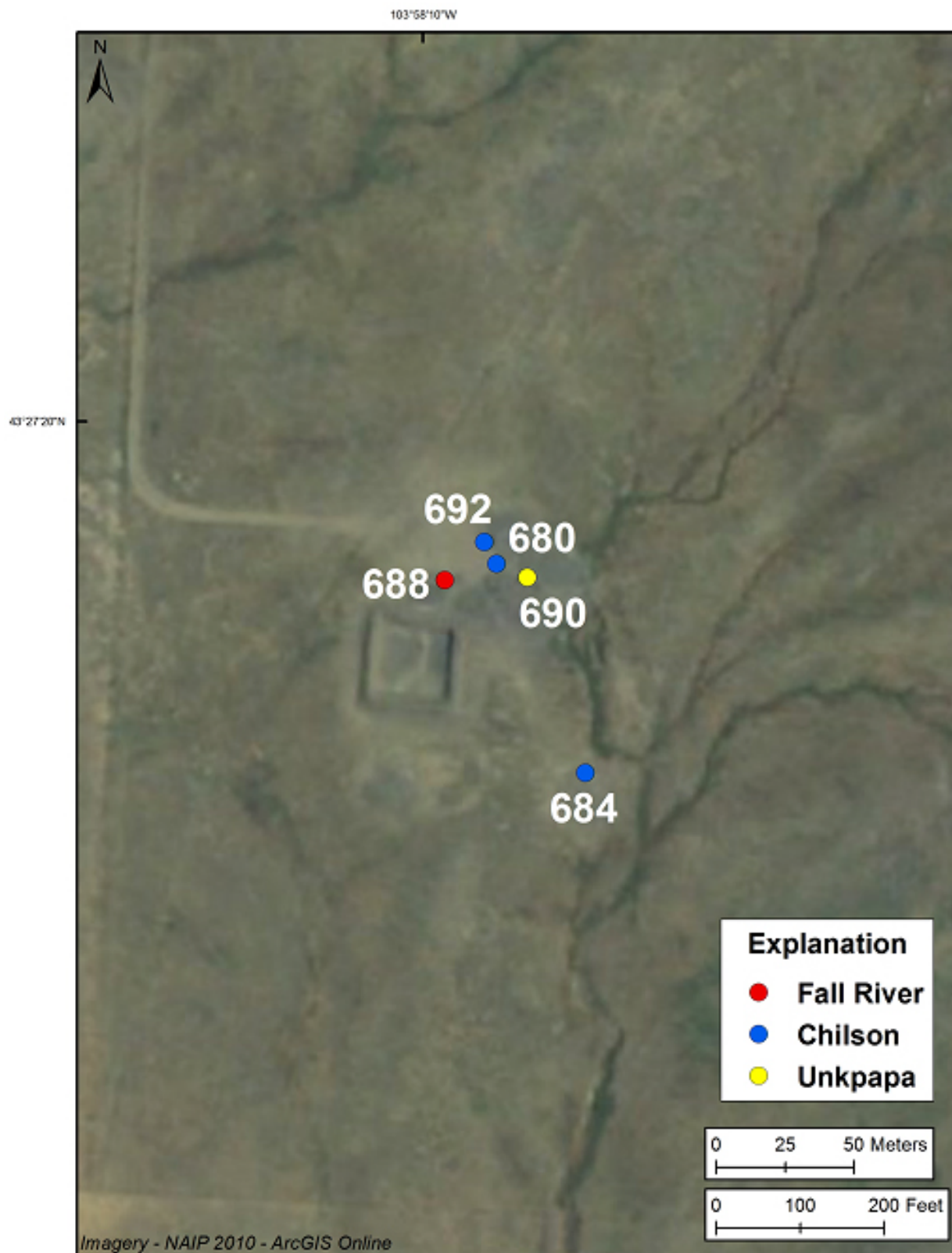


**Figure 2.** Satellite imagery overlain with monitoring well locations. Labels indicate well identification number. White boxes indicate location of expanded views for figures 3 and 4.





**Figure 3.** Expanded view of Dewey area. Labels indicate well identification number.

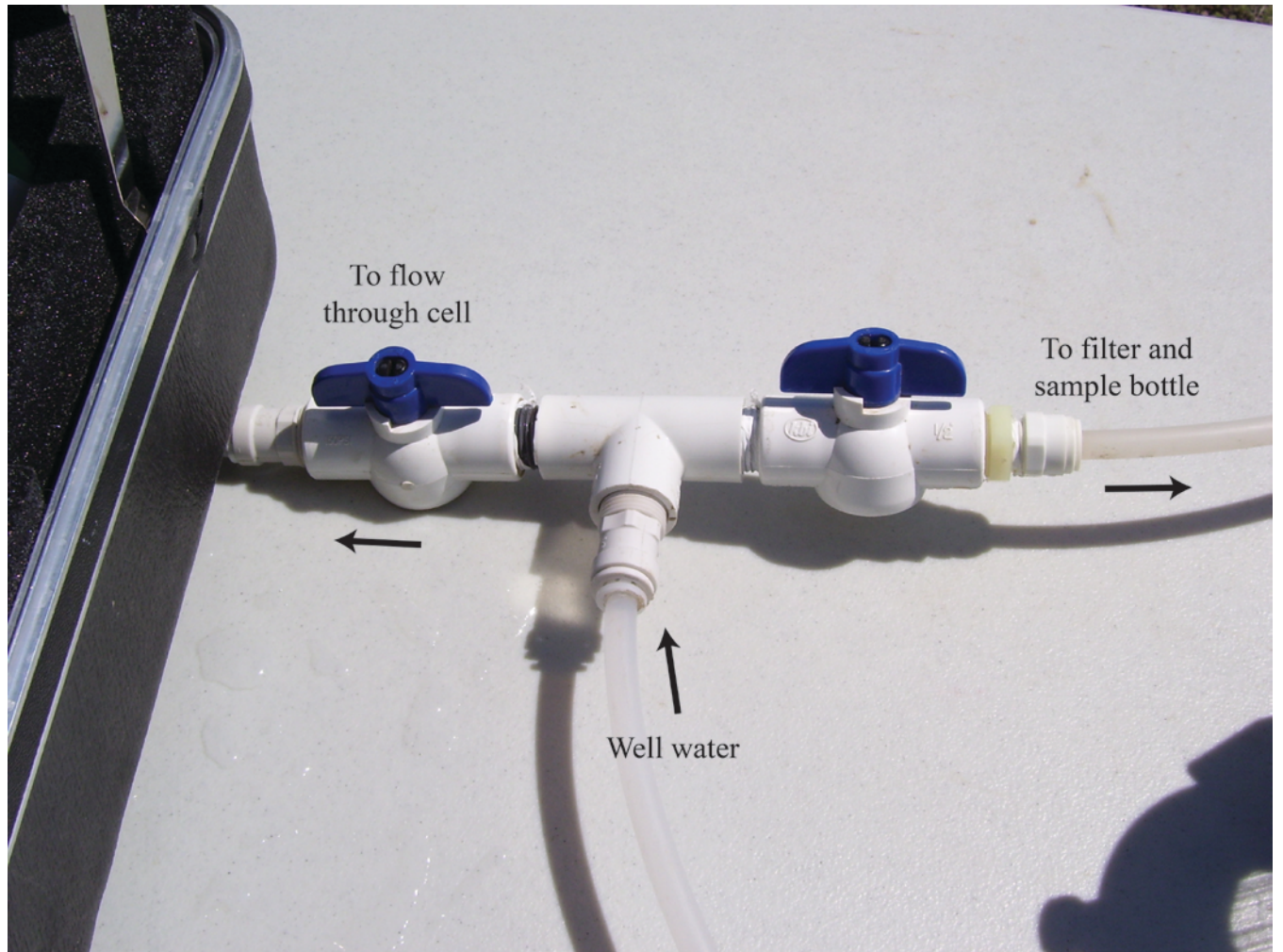


**Figure 4.** Expanded view of Burdock area. Labels indicate well identification number.





**Figure 5.** Typical wellhead control valves.



**Figure 6.** "T-valve" used for sampling.

[illegible]

Year	Month	Day	Time	Location	Activity	Remarks
2023	01	01	08:00	Room 101	Check-in	Guest from New York
2023	01	02	09:30	Room 102	Check-out	Guest from London
2023	01	03	10:15	Room 103	Check-in	Guest from Tokyo
2023	01	04	11:00	Room 104	Check-out	Guest from Paris
2023	01	05	12:30	Room 105	Check-in	Guest from Sydney
2023	01	06	13:45	Room 106	Check-out	Guest from Melbourne
2023	01	07	14:20	Room 107	Check-in	Guest from Auckland
2023	01	08	15:10	Room 108	Check-out	Guest from Christchurch
2023	01	09	16:00	Room 109	Check-in	Guest from Dunedin
2023	01	10	17:30	Room 110	Check-out	Guest from Invercargill
2023	01	11	18:15	Room 111	Check-in	Guest from Queenstown
2023	01	12	19:00	Room 112	Check-out	Guest from Milford Sound
2023	01	13	20:30	Room 113	Check-in	Guest from Fiordland
2023	01	14	21:15	Room 114	Check-out	Guest from Stewart Island
2023	01	15	22:00	Room 115	Check-in	Guest from Pukekohe
2023	01	16	23:45	Room 116	Check-out	Guest from Wairarapa
2023	01	17	00:30	Room 117	Check-in	Guest from Hawke's Bay
2023	01	18	01:15	Room 118	Check-out	Guest from Bay of Plenty
2023	01	19	02:00	Room 119	Check-in	Guest from Tairāwhiti
2023	01	20	02:45	Room 120	Check-out	Guest from Manawatu
2023	01	21	03:30	Room 121	Check-in	Guest from Capital
2023	01	22	04:15	Room 122	Check-out	Guest from North Island
2023	01	23	05:00	Room 123	Check-in	Guest from South Island
2023	01	24	05:45	Room 124	Check-out	Guest from West Coast
2023	01	25	06:30	Room 125	Check-in	Guest from East Coast
2023	01	26	07:15	Room 126	Check-out	Guest from Central
2023	01	27	08:00	Room 127	Check-in	Guest from North
2023	01	28	08:45	Room 128	Check-out	Guest from South
2023	01	29	09:30	Room 129	Check-in	Guest from West
2023	01	30	10:15	Room 130	Check-out	Guest from East
2023	01	31	11:00	Room 131	Check-in	Guest from Central
2023	02	01	11:45	Room 132	Check-out	Guest from North
2023	02	02	12:30	Room 133	Check-in	Guest from South
2023	02	03	13:15	Room 134	Check-out	Guest from West
2023	02	04	14:00	Room 135	Check-in	Guest from East
2023	02	05	14:45	Room 136	Check-out	Guest from Central
2023	02	06	15:30	Room 137	Check-in	Guest from North
2023	02	07	16:15	Room 138	Check-out	Guest from South
2023	02	08	17:00	Room 139	Check-in	Guest from West
2023	02	09	17:45	Room 140	Check-out	Guest from East
2023	02	10	18:30	Room 141	Check-in	Guest from Central
2023	02	11	19:15	Room 142	Check-out	Guest from North
2023	02	12	20:00	Room 143	Check-in	Guest from South
2023	02	13	20:45	Room 144	Check-out	Guest from West
2023	02	14	21:30	Room 145	Check-in	Guest from East
2023	02	15	22:15	Room 146	Check-out	Guest from Central
2023	02	16	23:00	Room 147	Check-in	Guest from North
2023	02	17	23:45	Room 148	Check-out	Guest from South
2023	02	18	00:30	Room 149	Check-in	Guest from West
2023	02	19	01:15	Room 150	Check-out	Guest from East
2023	02	20	02:00	Room 151	Check-in	Guest from Central
2023	02	21	02:45	Room 152	Check-out	Guest from North
2023	02	22	03:30	Room 153	Check-in	Guest from South
2023	02	23	04:15	Room 154	Check-out	Guest from West
2023	02	24	05:00	Room 155	Check-in	Guest from East
2023	02	25	05:45	Room 156	Check-out	Guest from Central
2023	02	26	06:30	Room 157	Check-in	Guest from North
2023	02	27	07:15	Room 158	Check-out	Guest from South
2023	02	28	08:00	Room 159	Check-in	Guest from West
2023	02	29	08:45	Room 160	Check-out	Guest from East
2023	02	30	09:30	Room 161	Check-in	Guest from Central
2023	03	01	10:15	Room 162	Check-out	Guest from North
2023	03	02	11:00	Room 163	Check-in	Guest from South
2023	03	03	11	Room 164	Check-out	Guest from West
2023	03	04	12	Room 165	Check-in	Guest from East
2023	03	05	13	Room 166	Check-out	Guest from Central
2023	03	06	14	Room 167	Check-in	Guest from North
2023	03	07	15	Room 168	Check-out	Guest from South
2023	03	08	16	Room 169	Check-in	Guest from West
2023	03	09	17	Room 170	Check-out	Guest from East
2023	03	10	18	Room 171	Check-in	Guest from Central
2023	03	11	19	Room 172	Check-out	Guest from North
2023	03	12	20	Room 173	Check-in	Guest from South
2023	03	13	21	Room 174	Check-out	Guest from West
2023	03	14	22	Room 175	Check-in	Guest from East
2023	03	15	23	Room 176	Check-out	Guest from Central
2023	03	16	00	Room 177	Check-in	Guest from North
2023	03	17	01	Room 178	Check-out	Guest from South
2023	03	18	02	Room 179	Check-in	Guest from West
2023	03	19	03	Room 180	Check-out	Guest from East
2023	03	20	04	Room 181	Check-in	Guest from Central
2023	03	21	05	Room 182	Check-out	Guest from North
2023	03	22	06	Room 183	Check-in	Guest from South
2023	03	23	07	Room 184	Check-out	Guest from West
2023	03	24	08	Room 185	Check-in	Guest from East
2023	03	25	09	Room 186	Check-out	Guest from Central
2023	03	26	10	Room 187	Check-in	Guest from North
2023	03	27	11	Room 188	Check-out	Guest from South
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2023	03	29	13	Room 190	Check-out	Guest from East
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2023	04	02	17	Room 194	Check-out	Guest from West
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2023	05	14	28	Room 237	Check-in	Guest from North
2023	05	15	29	Room 238	Check-out	Guest from South
2023	05	16	30	Room 239	Check-in	Guest from West
2023	05	17	31	Room 240	Check-out	Guest from East
2023	05	18	00	Room 241	Check-in	Guest from Central
2023	05	19	01	Room 242	Check-out	Guest from North
2023	05	20	02	Room 243	Check-in	Guest from South
2023	05	21	03	Room 244	Check-out	Guest from West
2023	05	22	04	Room 245	Check-in	Guest from East
2023	05	23	05	Room 246	Check-out	Guest from Central
2023	05	24	06	Room 247	Check-in	Guest from North
2023	05	25	07	Room 248	Check-out	Guest from South
2023	05	26	08	Room 249	Check-in	Guest from West
2023	05	27	09	Room 250	Check-out	Guest from East
2023	05	28	10	Room 251	Check-in	Guest from Central
2023	05	29	11	Room 252	Check-out	Guest from North
2023	05	30	12	Room 253	Check-in	Guest from South
2023	05	31	13	Room 254	Check-out	Guest from West
2023	06	01	14	Room 255	Check-in	Guest from East
2023	06	02	15	Room 256	Check-out	Guest from Central
2023	06	03	16	Room 257	Check-in	Guest from North
2023	06	04	17	Room 258	Check-out	Guest from South
2023	06	05	18	Room 259	Check-in	Guest from West
2023	06	06	19	Room 260	Check-out	Guest from East
2023	06	07	20	Room 261	Check-in	Guest from Central
2023	06	08	21	Room 262	Check-out	Guest from North
2023	06	09	22	Room 263	Check-in	Guest from South
2023	06	10	23	Room 264	Check-out	Guest from West
2023	06	11	24	Room 265	Check-in	Guest from East
2023	06	12	25	Room 266	Check-out	Guest from Central
2023	06	13	26	Room 267	Check-in	Guest from North
2023	06	14	27	Room 268	Check-out	Guest from South
2023	06	15	28	Room 269	Check-in	Guest from West
2023	06	16	29	Room 270	Check-out	Guest from East
2023	06	17	30	Room 271	Check-in	Guest from Central
2023	06	18	31	Room 272	Check-out	Guest from North
2023	06	19	00	Room 273	Check-in	Guest from South
2023	06	20	01	Room 274	Check-out	Guest from West
2023	06	21	02	Room 275	Check-in	Guest from East
2023	06	22	03	Room 276	Check-out	Guest from Central
2023	06	23	04	Room 277	Check-in	Guest from North
2023	06	24	05	Room 278	Check-out	Guest from South
2023	06	25	06	Room 279	Check-in	Guest from West
2023	06	26	07	Room 280	Check-out	Guest from East
2023	06	27	08	Room 281	Check-in	Guest from Central
2023	06	28	09	Room 282	Check-out	Guest from North
2023	06	29	10	Room 283	Check-in	Guest from South
2023	06	30	11	Room 284	Check-out	Guest from West
2023	06	31	12	Room 285	Check-in	Guest from East
2023	07	01	13	Room 286	Check-out	Guest from Central
2023	07	02	14	Room 287	Check-in	Guest from North
2023	07	03	15	Room 288	Check-out	Guest from South
2023	07	04	16	Room 289	Check-in	Guest from West
2023	07	05	17	Room 290	Check-out	Guest from East
2023	07	06	18	Room 291	Check-in	Guest from Central
2023	07	07	19	Room 292	Check-out	Guest from North
2023	07	08	20	Room 293	Check-in	Guest from South
2023	07	09	21	Room 294	Check-out	Guest from West
2023	07	10	22	Room 295	Check-in	Guest from East
2023	07	11	23	Room 296	Check-out	Guest from Central
2023	07	12	24	Room 297	Check-in	Guest from North
2023	07	13	25	Room 298	Check-out	Guest from South
2023	07	14	26	Room 299	Check-in	Guest from West
2023	07	15	27	Room 300	Check-out	Guest from East
2023	07	16	28	Room 301	Check-in	Guest from Central
2023	07	17	29	Room 302	Check-out	Guest from North
2023	07	18	30	Room 303	Check-in	Guest from South
2023	07	19	31	Room 304	Check-out	Guest from West
2023	07	20	00	Room 305	Check-in	Guest from East
2023	07					