1 Swiss data quality: augmenting CAMELS-CH with isotopes, water

2 quality, agricultural and atmospheric data

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21 Abstract

22 Despite the growth of large-sample hydrology (LSH) datasets, water quality data remain scarce. Here, we introduce CAMELS-CH-Chem, an extension of CAMELS-CH (Catchment 23 24 Attributes and Meteorology for Large-sample Studies in Switzerland), incorporating up to 40 25 water guality parameters for 115 Swiss catchments from 1981 to 2020. The dataset includes 26 hourly and daily time series of measurements of water temperature, dissolved oxygen, pH, 27 and electrical conductivity; as well as (bi)monthly measurements of alkalinity (HCO₃₎, 28 ammonium, Ca, Cl, dissolved organic carbon (DOC), dissolved reactive phosphorus (DRP), total 29 organic carbon (TOC), K, Mg, Na, total filtered phosphorus, total hardness, total nitrogen, 30 total phosphorus, NO₃, NO₂, Si, SO₄, and stream isotopes. In addition, we provide catchment 31 aggregated (bi)monthly time series of rainwater isotopes, along with annual data on land 32 cover, agricultural data (crop types and livestock density), and atmospheric nitrogen 33 deposition. This comprehensive dataset enables broader integration of water quality into LSH 34 research and supports new insights specifically in the field of hydrological modelling.

35 Background & Summary

Recently, there has been a widespread development of large-sample hydrology (LSH) datasets worldwide¹⁻¹⁰. Many of these datasets were inspired by the pioneering Catchment Attributes and MEteorology for Large-sample Studies (CAMELS) dataset, which provided a comprehensive LSH dataset for the contiguous United States⁶. Such datasets typically include hydro-climatic variables—such as streamflow, meteorological forcing data, and catchment properties (e.g., land use and soil types)—covering numerous catchments over extended time periods.

Yet, numerous studies have highlighted the importance of integrating long-term hydroclimatic and catchment properties with stream water quality data to derive critical insights into solute transport processes^{11,12}. Hence, datasets that combine long-term, reliable water quality variables with other hydro-climatic and catchment properties are essential to address 47 global environmental challenges arising from population growth, land use intensification and48 a changing climate.

Recently, CAMELS-Chem¹¹ was released as an openly accessible dataset for the contiguous United States—the first augmentation of a CAMELS dataset that incorporates water quality data. However, similar initiatives remain limited and freely accessible water quality parameters remain scarce in published datasets. This is primarily due to the challenges associated with measuring and providing access to such data, compared to hydro-climatic and catchment variables that are typically easier to obtain.

55 Here, we introduce CAMELS-CH-Chem, an extension of the existing CAMELS-CH⁵ dataset. 56 While CAMELS-CH provides hydrometeorological and streamflow data for 331 catchments 57 across Switzerland, CAMELS-CH-Chem builds on this by integrating up to 40 aggregated stream 58 water quality parameters and isotopes. It also includes catchment-aggregated data on 59 atmospheric deposition, land cover, and agriculture for a subset of 115 catchments. With 60 CAMELS-CH-Chem, we aim to make a significant contribution to the field of LSH by introducing 61 the first CAMELS extension in Europe to include water quality data, enabling data-driven 62 modeling of water quality at larger scales.

63 Our intention in providing catchment-aggregated variables is to enable comprehensive, multi-64 scale analyses by offering a harmonized dataset ready for integration into LSH and hydro 65 chemical modelling frameworks. Catchment aggregated data of agricultural practices and 66 atmospheric deposition serve as critical explanatory variables for interpreting spatial and 67 temporal variability in water quality and biogeochemical parameters. Similarly, the inclusion 68 of precipitation isotope data adds valuable insights into hydrological flow paths, water source 69 contributions, and catchment-scale transit times, which can be used alongside stream water 70 isotope time series.

Although some of the original data, dating back to 1970, is available upon request from providers such as FOEN¹³, CAMELS-CH-Chem specifically provides a comprehensive dataset from 1981 to 2020 to maximize the overlap between different water quality sources and the complementary CAMELS-CH. This approach aligns with the primary objective of LSH datasets: to provide long-term, standardized variables across large regions¹⁴.

76 The newly provided data is divided into three main categories:

- 77 Stream water chemistry: This includes time series of more than 30 stream water (i) 78 chemistry constituents, covering both field and laboratory measurements. It provides 79 hourly and daily data on water temperature, dissolved oxygen, pH, and electrical 80 conductivity, along with (bi)monthly measurements of alkalinity, ammonium (NH_4^+), 81 calcium (Ca), chloride (Cl), dissolved organic carbon (DOC), dissolved reactive 82 phosphorus (DRP), total organic carbon (TOC), bicarbonate (HCO₃⁻), potassium (K), 83 magnesium (Mg), sodium (Na), total hardness, total dissolved nitrogen (TDN), total 84 organic nitrogen (TON), total phosphorus (TP), nitrate (NO₃[−]), nitrite (NO₂[−]), silica (Si), and sulphate (SO_4^{2-}) . 85
- 86 (ii) Stream water isotopes: This includes (bi)monthly time series of stream water isotope
 87 data of deuterium (²H) and oxygen-18 (¹⁸O).
- 88 (iii) **Catchment aggregated data:** This provides annual time series of atmospheric 89 deposition concentrations for nitrate (NO_3^-) , ammonium (NH_4^+) , ammonia (NH_3) , 90 nitrite (NO_2^-) , and total inorganic nitrogen for 115 catchments, alongside crop type 91 distributions and livestock density data. Finally, we also provide monthly time series 92 of catchment aggregated rainwater isotope data to improve isotope assessments.

93 The manuscript is structured as follows: the Methods section describes the original data 94 sources and the methodology used for compiling it into the CAMELS-CH-Chem dataset. The 95 Data Records section describes the structure of the CAMELS-CH-Chem dataset. Finally, the 96 Technical Validation section provides a first order validation of the CAMELS-CH-Chem dataset 97 based on selected hypotheses.

98 Methods

99 Stream water chemistry data

100 Water chemistry data was collected within the framework of national monitoring programmes. The continuous water temperature data was obtained from the surface water 101 102 temperature monitoring network of the Swiss Federal Office for the Environment (FOEN, in 103 German BAFU). The water quality data was collected within the National Surface Water 104 Quality Monitoring Programme (NAWA). Within NAWA TREND, the surface water quality is 105 monitored in more than 100 catchments in cooperation with the FOEN and the cantonal 106 authorities. In NAWA FRACHT (previously called NADUF), the continuous parameters and 107 pollutant loads are monitored in about 15 selected catchments in collaboration with the FOEN, 108 the Swiss Federal Institute of Aquatic Science and Technology (Eawag) and the Swiss Federal 109 Institute for Forest, Snow and Landscape Research (WSL). The three sources encompass both 110 unique and redundant information for a few variables, whereby continuous (sensors) data 111 comprises the most complete daily and hourly time series.

112 Figure 1 shows the distribution of the stream water chemistry measurement stations across 113 Switzerland with their respective catchment boundaries in background. The colors of each dot 114 represent the maximum number of chemical variables observed at a given location. Note that some locations of the three different data sources overlap, consequently we provide data for 115 116 115 unique locations. Figure 1 also illustrates that these catchments are well distributed across 117 Switzerland capturing the complex topography (Figure 1d) and climatology of the country. In 118 total, 86 locations have available continuous measurement data, 24 have available water 119 sampling data from NAWA FRACHT and 76 from NAWA TREND.



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Figure 1. Spatial distribution of the measurement locations with data available in CAMELS-CH-Chem encompassing (a) Sensor data, (b) NAWA FRACHT and (c) NAWA TREND data. The dots in each subplot have different colors representing the number of available parameters at each location. Note that for the same location, we might provide data from continuous measurements, NAWA FRACHT and NAWA TREND. The upstream catchment area of each station is displayed in background. Additionally, subplot (d) shows a map of the elevation, with the boundaries of Switzerland and the catchment areas of each station.

128 In the following section, we describe the main data sources. A detailed description of the 129 measurements and information on data acquisition and processing, such as sensor types, 130 accuracy, and methods used, are available in Supplementary Material.

131 Continuous (sensors) data

The FOEN¹³ and NAWA FRACHT¹⁵ programme provided hourly and daily time series of stream water temperature, pH, electric conductivity, and oxygen concentration from 1981 to 2020 for the 86 stations shown in **Figure 1a**. It is important to note that most of the 86 stations have temperature only. These data are continuously measured with online sensors. Here we refer to them as "continuous data". A further overview of these four variables regarding the datasetspecific information (name and description), units and resolution is shown in **Table 1**.

Table 1. Overview of the stream water chemistry data obtained from the continuous dataprovided by FOEN and the NAWA FRACHT programme.

Attribute	Description	Units	Temporal resolution	Source
date	Date of the measurement.	-	Hourly, daily	FOEN ¹³ and
temp_sensor	Water temperature	°C		FRACHT ¹⁵

Attribute	Description	Units	Temporal resolution	Source
ec_sensor	Electrical conductivity at 25 °C	μS/cm		
O2C_sensor	Oxygen concentration	mg/l		
pH_sensor	рН	-		

140 NAWA FRACHT

Further water chemistry data was obtained from the NAWA FRACHT programme¹⁵ (Figure 1b). 141 The dataset provides 38 variables obtained from either installed online sensors (six variables 142 with "_sensor" in their names) or measured in the laboratory (remaining variables) (Table 2). 143 144 This data has a measurement resolution of 7 to 14 days collected between 1982 and 2020, 145 whereby time series provide the mean of measurements between date_start and date_end. 146 Note that the NAWA FRACHT program also has overlapping measurement locations with the 147 continuous data (described above) and NAWA TREND (described below). An overview of the 148 38 measured variables, i.e., dataset-specific information, units and resolution, is shown in 149 Table 2.

150 **Table 2.** Overview of the stream water chemistry variables obtained from the National River

151 Monitoring and Survey Programme (NAWA FRACHT, previously called NADUF). The time series

152 provide the mean of measurements between date_start and date_end.

Attribute	Description	Units	Temporal	Source
			resolution	
date_start	Measurement start date.	-		
date_end	Measurement end date.	-		
alk	Alkalinity	mmol/l		
As	Arsenic	μg/l		
Ва	Barium	μg/I		
Br	Bromide	mg/l		
Cd	Cadmium	μg/l		
Са	Calcium	mg/l		
Cl	Chloride	mg/l		
Cr	Chromium	μg/I		
Cu	Copper	μg/l	7 14 dove	NAWA
doc	Dissolved Organic Carbon	mg/l		
drp	Dissolved Reactive Phosphorus	mg/l	7-14 Udys	FRACHT
ec25_sensor*	Electrical conductivity at 25 °C	μS/cm	mean	15
ec20_lab	Electrical conductivity at 20 °C	μS/cm		
F	Fluoride	mg/l		
Fe	Iron	mg/l		
Pb	Lead	μg/l		
Mg	Magnesium	mg/l		
q_mean_sensor*	Mean discharge	m³/s		
Hg	Mercury	μg/l		
Ni	Nickel	μg/I		
NO3_N	Nitrate	mg/l		
O2C_sensor*	Oxygen concentration	mg/l		
O2S_sensor*	Oxygen saturation	%		

Attribute	Description	Units	Temporal resolution	Source
pH lab	рН	-		
pH_sensor*	рН	-		
К	Potassium	mg/l		
H4SiO4	Silicate	mg/l		
Na	Sodium	mg/l		
Sr	Strontium	μg/l		
SO4	Sulphate	mg/l		
tfp	Total filtered phosphorus	mg/l		
th	Total hardness	mmol/		
tn	Total nitrogen	mg/l		
toc	Total organic carbon	mg/l		
tp	Total phosphorus	mg/l		
tss	Total suspended solids	mg/l		
temp_sensor*	Water temperature	°C		
Zn	Zinc	μg/l		

153 * These values are averages computed for the **date_start** and **date_end** measurement interval

and derived from sensors installed at the measurement location with an original resolution of

155 10-minutes (FOEN and NAWA FRACHT).

156 NAWA TREND

Water chemistry data is also provided from the NAWA TREND programme¹⁶ (**Figure 1c**). This dataset provides 22 variables (**Table 3**), measured from grab samples covering 2011 through 2020 at monthly resolution. Thus, the time series represents the measurement taken at the respective **date**. An overview of the 22 variables, including dataset-specific information, units,

161 and resolution, is shown in **Table 3**.

Table 3. Overview of the stream water chemistry variables obtained from the National Surface
 Water Quality Monitoring Programme (NAWA TREND). The measurements were taken as grab

164 samples, typically once per month. The precise sampling dates are provided in the **date** 165 column in the final dataset.

Attribute	Description	Units	Temporal	Source
			resolution	
date	Date the measurement was taken.	-		
NH4_N	Ammonium nitrogen	mg/l		
Cl	Chloride	mg/l		
q_max_kanton	Daily maximum discharge	m³/s		
	measured or estimated from		One grab	
	cantons			
q_min_kanton	Daily minimum discharge measured	m³/s	sample per	
	or estimated from cantons		monun,	
q_mean_kanton	Daily mean discharge measured or	m³/s	Detween	IRENU
	estimated from cantons		2011 anu	
q_mean_sensor*	Mean discharge	m³/s	2020	
doc	Dissolved organic carbon	mg/l		
drp	Dissolved reactive phosphorus	mg/l		
ec25_lab	Electrical conductivity at 25 °C	μS/cm		
	measured in the lab			

ec25_sensor*	Electrical conductivity at 25 °C	μS/cm	
NO3_N	Nitrate nitrogen	mg/l	
NO2_N	Nitrite nitrogen	mg/l	
O2_lab	Oxygen concentration measured in the lab	mg/l	
O2_sensor*	Oxygen concentration	mg/l	
O2S _sensor*	Oxygen saturation	%	
pH_lab	pH measured in the lab	-	
pH_sensor*	рН	-	
temp_lab	Water temperature measured in the lab	°C	
temp_sensor*	Water temperature	°C	
turbidity_sensor*	Turbidity	NTU	
tn	Total nitrogen	mg/l	
tp	Total phosphorus	mg/l	

167 * These values are averages computed from sensors installed at the measurement location at

168 a resolution of 10-minutes (FOEN¹³ and NAWA FRACHT¹⁵) for the respective measurement 169 date.

170 Stream water isotopes

We also provide stream water measurements of deuterium and oxygen-18 data with a resolution from 14 days to monthly for all locations where such data is available across Switzerland from the ISOT module¹⁷ (nine stations) of the National Groundwater Monitoring (NAQUA) and from the CH-IRP dataset¹⁸ (11 stations). The spatial distribution of such stations is shown in **Figure 2**.



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Figure 2. Spatial distribution of the measurement locations with stream water isotope data available in CAMELS-CH-Chem encompassing (a) ISOT data and (b) CH-IRP. Their respective catchment delineations are shown in background for both datasets. Moreover, for CH-IRP, it is worth noting that stations Biberbrugg (2604) and Einsiedeln (2609), both depicted as red dots, are located very close but in two different rivers (Biber and Alp). Therefore, due to scale reasons this figure gives the impression of having only 10 stations.

184 ISOT module

The ISOT module provides isotopes measurement time series covering the period from 1992 through 2022. For two stations (i.e., Thun: 2030 & Brienzwiler: 2019), the measurements are from grab samples taken at the respective time or a mixed sample for multiple timesteps of the same month in case multiple dates are provided. Information on the data is summarized in (**Table 4**). Finally, **Figure 2a** shows the distribution of these nine measurement locations. 190 **Table 4.** Overview of the ISOT isotope data available in CAMELS-CH-Chem. The resolution is

191 variable for each time-step, and ranges from 14 days to monthly. The provided measurements

are the average from daily samples between **date_start** and **date_end**, whereas aggregation

193 intervals vary up to a maximum of 30 days.

Attribute	Description	Units	Temporal resolution	Source
date_start	Measurement start date.	-		
date_end	Measurement end date.	-	Monthly or 14-day average	ISOT ¹⁷
delta_2h	Deuterium (² H)	δ^{2} H ‰ SMOW		
delta_18o	Oxygen-18 (¹⁸ O)	δ^{18} O ‰ SMOW		

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195CH-IRP dataset

196 Apart from the nine stations monitored by the ISOT module, here we also provide data for 11 197 stations monitored and made available by the CH-IRP dataset and project (Staudinger, 2019¹⁸), 198 covering the period from 2010 through 2020. The CH-IRP original dataset covers a total of 22 199 medium-sized alpine and pre-alpine Swiss catchments. It is worth noting that we deliberately 200 provide data only for the 11 stations overlapping with the original CAMELS-CH stations. For a 201 full description of the CH-IRP dataset, users should refer to their original publication (Staudinger, 2019¹⁸). Information on the data is summarized in (Table 5), while Figure 2b 202 203 shows the distribution of these monitoring stations.

Table 5. Overview of the CH-IRP isotope data available in CAMELS-CH-Chem. This data is
 measured every 14 days. The date attribute corresponds to the sampling date.

Attribute	Description	Units	Temporal	Source
			resolution	
date	Measurement	-		
	date.		F 44 1	CI II 2010 ¹⁸
delta_2h	Deuterium (² H)	δ^{2} H ‰ SMOW	Every 14-days	Staudinger, 2019 ¹⁰
delta_18o	Oxygen-18 (¹⁸ O)	δ^{18} O ‰ SMOW		

206 Catchment aggregated data

207 Complementing the stream water chemistry and isotope data, CAMELS-CH-Chem also 208 provides five types of catchments aggregated data: yearly time series of i) atmospheric 209 deposition, ii) land cover percentage, iii) crop types and iv) livestock unit data, along with (v) 210 monthly time series of rainwater isotopes data. **Figure 3** shows the locations of the 115 211 catchments across Switzerland where these aggregated data is available. Note that these 212 catchments correspond to those with stream water chemistry data.

It is important to note that, except for land cover data, this aggregated information was derived from data sources covering solely Switzerland. Since 23 of the 115 catchments have a part located outside Switzerland, users should be careful when dealing with the data. To provide an initial filter for users, the variable **area_perc_swiss**, which will be further described in the Gauge metadata section (**Table 11**), provides the percentage of the catchment located in Switzerland.



Figure 3. Spatial distribution of the 115 catchments used to derive the catchment aggregated data provided in CAMELS-CH-Chem. Each of their respective gauging stations is shown as black crosses. In the background, the four major basins are shown, along with their main river networks and major lakes.

224 Atmospheric deposition

225 CAMELS-CH-Chem also provides time series of annual atmospheric deposition of NO₃⁻, NH₄⁻, NH₃, NO₂, HNO₃, and total inorganic nitrogen aggregated from gridded data provided by Rihm 226 227 and Künzle, 2019¹⁹. Specifically, the gridded data of individual nitrogen components was based 228 on i) emission inventories and statistical dispersion models (NH_3 and NO_2), ii) monitoring data and spatial interpolation methods (HNO₃, wet deposition of NO₃⁻ and NH₄⁻), and iii) monitoring 229 230 data, inferential models, and spatial interpolation (dry deposition of NO₃⁻ and NH₄⁻). Finally, the 231 gridded data of total atmospheric nitrogen deposition is based on the combination of the 232 above-mentioned components.

Further details on the methods used to model and spatially aggregate nitrogen deposition are available in Rihm and Achermann, 2016²⁰ and Rihm and Künzle, 2019¹⁹. The original gridded nitrogen deposition data is available at a 1 km x 1 km resolution in 5-year intervals starting in 1990. Here, we provide catchment averages, which were calculated using the area-weighted mean of all map pixels inside a catchment. **Table 6** provides an overview of this dataset. It is worth noting that since this data exists only for the years 1990, 2000, 2005, 2010, 2015 and 2020, we applied a linear interpolation between these to fill the years in between.

J	Table 6. Overview of the atmospheric deposition data available in CAMELS-CH-Chem.						
	Attribute	Description		Units	Temporal	Source	
					resolution		
	date	Year of	the	-		19,21	
		incusurement.					

240 **Table 6.** Overview of the atmospheric deposition data available in CAMELS-CH-Chem.

Attribute	Description	Units	Temporal resolution	Source
dhno3gas	Gaseous deposition of HNO ₃			
dnh3gas	Gaseous deposition of NH ₃			
dnh4total	Sum of wet and dry deposition of NH ₄ ⁻		Voorly	
dno2gas	Gaseous deposition of NO ₂ ⁻	kg N/ ha	between 1990	
dno3total	Sum of wet and dry deposition of NO ₃ -			
dntotal	Total nitrogen deposition as a sum of wet, dry and gaseous deposition			

241 Landcover data

242 We provide land use coverage for the 115 catchments included in CAMELS-CH-Chem, recomputed following the same procedure as in CAMELS-CH⁵, also using the CORINE Land 243 Cover (CLC) dataset²². As CLC data is available for the years 2000, 2006, 2012 and 2018, we 244 245 applied linear interpolation to fill the years in between and repeated the value 2018 for the 246 last two years. The data was divided into 12 classes: agriculture, forest (coniferous, deciduous, 247 and mixed), grass and herb vegetation, scrub vegetation, wetlands, ice and perpetual snow, inland water surface, rock (loose and solid), settlements/urban and unknown/blank. Table 7 248 249 summarizes this information. Users may refer to CAMELS-CH⁵ and the official CORINE²² 250 publication for further details.

251 **Table 7.** Overview of the land cover data provided in CAMELS-CH-Chem.

Attribute	Description	Units	Temporal	Source
			resolution	
date	Year of the measurement.	-		
crop_perc	Agriculture			
dwood_perc	Deciduous forest			
ewood_perc	Coniferous forest (evergreen)			
grass_perc	Grass and herb vegetation			
ice_perc	Glaciers and perpetual snow		Yearly,	
inwater_perc	Inland water	%	2000 and	CORINE ²²
loose_rock_perc	Loose rocks and bare soils		2000 and 2020	
mixed_wood_per c	Mixed forest		2020	
rock_perc	Hard rocks and bare soils			
scrub_perc	Percentage of medium- scale vegetation			
urban_perc	Urban and settlements			
wetlands_perc	Wetlands			

252 Crop types data

We also estimated the area within the 115 catchments covered by certain crops and provided these data annually from 1980 to 2020. The following 10 crop types were considered: cereals, maize, sugar beet, potatoes, rapeseed, pulses, vegetables, total arable land (= sum of all crops), as well as grapevines and orchards. We utilized the Swiss census of agricultural farms, provided by FSO, 2023²³, for the annual statistics of all crops in Switzerland.

258 Until 2019, yearly crop statistics were recorded only at the municipal level, meaning the 259 precise location of crops within each municipality was unknown. Hence, to improve spatial 260 localization, we distributed the statistical crop data by aggregating such yearly values across 261 the land use class 41 (period 2004-09, and standard nomenclature NOAS04) obtained from the Arealstatistik Schweiz dataset²⁴ for the arable land; and the classes "grapevine" and "orchard" 262 from the Topographic Landscape Model (TLM) from the Swisstopo²⁵ dataset. This step 263 264 resulted in the total crop data being divided into 10 different classes for each Swiss 265 municipality.

Finally, we aggregated the municipality data per catchment and estimated the area of each crop type for each of the 115 catchments. Each catchment has, therefore, a yearly time series for each of the 10 crop classes (**Table 8**). It is important to note that the data before 1996 was provided at 5-year intervals (1980, 1985, 1990 and 1996), and after 1996, at yearly time-steps until 2019. Therefore, we applied a linear interpolation between the 5-year data in the 1980– 1996 period and repeated the values from 2019 for the last year.

Table 8. Overview of the crop type data available in CAMELS-CH-Chem, along with theirrespective temporal resolution and source.

Attribute		Units	Temporal resolution	Source
date	Year of the measurement.	-		
	cereal	-		
	maize		Yearly, between 1980 and 2020	
Anglela land	sugarbeet			
	potato			
Alable latiu	rapeseed			23–25
	pulse	111-		
	vegetable			
	total_arable			
grapevine]		
	orchard			

274 Livestock unit data

The term livestock unit (here referred to as GVE, from the German word *Grossvieheinheiten*) is a reference unit that facilitates the aggregation of livestock across different species and age groups based on a standardized convention from Eurostat²⁶. Here, we used yearly livestock unit data from the Swiss Federal Statistical Office (FSO, 2023²⁴), covering the years 1980 to 2020. The original data was recorded at the municipal level, meaning the exact location of livestock within a municipality could not be determined.

Therefore, to improve spatial localization, we distributed the livestock data across the land use categories alpine and jura pastures, natural meadows, and farm pastures within each municipality. This was done by using land use classifications from the *Arealstatistik Schweiz* dataset (FSO, 2024²⁷), allowing us to estimate livestock density (livestock units per hectare)
 for different land use types, including natural meadows, pastures, and Alpine and Jura
 pastures.

- 287 We distinguished between two types of areas:
- Alpine and Jura Pastures: It is estimated that 20% of the total Swiss livestock population spends three months annually on these pastures. To calculate livestock density in these areas, we multiplied the total Swiss livestock units by 0.05 (20% × 1/4 year). This value was then divided by the total area of Alpine and Jura pastures, resulting in a uniform livestock unit per hectare for all such pastures.
- 293
 2. Natural Meadows and Farm Pastures: We used land use categories 15 (natural meadows) and 16 (home pastures) from the Swiss land use statistics²⁴. Each area was assigned a weighting factor of 1 and multiplied by the total livestock unit of the respective municipality. The resulting value was then multiplied by 0.95 (i.e., 1-0.05) and divided by the total area of natural meadows and home pastures within the municipality, yielding a municipality-specific livestock unit per hectare.

Finally, we aggregated the livestock data for the 115 catchments. An overview of the finallivestock unit data is presented in Table 9.

301 It is important to note that similarly to the crop-types data, livestock unit data before 1996 302 were provided at 5-year intervals (1980, 1985, 1990 and 1996) and after 1996, at yearly time-303 step until 2020. Therefore, we applied a linear interpolation between the years 1980 and 1996.

Table 9. Overview of the livestock unit data (GVE) available in CAMELS-CH-Chem, alongside their description, units, temporal resolution, and sources. Note that from 1980 to 1990, data is provided every five years, and from 1996 to 2022, data is provided yearly.

	- /	,		/
Attribute	Description	Units	Temporal	Source
			resolution	
date	Year of the	-		
	measurement.			
gve_sum	Number of	unit		
	livestock units		Yearly, between	24.26.27
	per catchment.		1980 and 2020	, -,
gve_ha	Number of	unit/ha		
	livestock units			
	per hectare.			

307 Rainwater isotopes data

Stable isotopes of oxygen (¹⁸O) and deuterium (²H) in precipitation and in stream water serve as natural tracers of hydrological processes. Hence, besides the stream water isotopes time series (previously described), we provide monthly catchment-aggregated rainwater isotopes for the 115 catchments from 2007 to 2020.

In Switzerland, stable isotopes of oxygen in precipitation are monitored through the ISOT¹⁷ observation network, which is part of the NAQUA National Groundwater Monitoring Programme. Here, we used monthly precipitation isotope values from the ISOT network²⁸, which were originally spatially interpolated into gridded isotope maps ("isoscapes") using a regression-kriging approach²⁹.

According to the isoscapes publication²⁸, this interpolation method involves a multiple linear regression model relating isotope values to a set of geographic and climatic variables, including

- 319 elevation, coordinates, and monthly precipitation totals. The spatially correlated residuals
- 320 from this regression are then interpolated using ordinary kriging to account for local deviations
- 321 not explained by the predictors.
- Table 10. Overview of the isoscapes rainwater isotope data available in CAMELS-CH-Chem.
 The data is provided at a monthly resolution. The date attribute is the sampling date.

Attribute	Description	Units	Temporal	Source
			resolution	
date	Measurement	-	Monthly	
	date.		between the	
delta_2h	Deuterium (² H)	δ^{2} H ‰ SMOW	years 2007 and	Isoscapes ²⁸
delta_18o	Oxygen-18 (¹⁸ O)	δ^{18} O ‰ SMOW	2020	

324 Catchment delineation

We used the catchment boundary shapefiles from the CAMELS-CH dataset to calculate 325 326 catchment aggregated data (i.e., atmospheric deposition and agricultural data). Catchment 327 outlets in CAMELS-CH are defined based on the discharge gauging location. However, some 328 chemical measurement locations in the NAWA FRACHT and NAWA TREND datasets are slightly 329 different from the CAMELS-CH streamflow gauging stations. For these cases, we adjusted the 330 CAMELS-CH catchment areas using the new outlet information. Information regarding these 331 shifts is provided in the gauge metadata (Table 11) with details regarding the distance 332 between streamflow and the water chemistry measurement locations. For the respective 333 catchments, we also provide the adjusted shapefile delineation for users to decide whether to 334 use the original CAMELS-CH or the adjusted CAMELS-CH-Chem catchment boundaries. All the 335 remaining catchment aggregated data were derived exclusively using the catchment 336 boundaries provided by CAMELS-CH.

337 Data Records

The current version of the CAMELS-CH-Chem dataset (v0.1)³⁰ is stored in a Zenodo repository
 at https://doi.org/10.5281/zenodo.14980027. The repository is organized into the following
 (sub)folders:

- *catchment_aggregated_data:* contains five subfolders. Each contains one csv file per catchment, with 115 files in total. The files are organized by time series (rows) and attribute variables (columns).
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- *agricultural_data*: contains one csv file per catchment with the variables described in **Table 8**.
- atmospheric_deposition: contains one csv file per catchment with the variables described in Table 6.
- *landcover_data:* contains one csv file per catchment with the variables described in Table 7.
- *livestock_data*: contains one csv file per catchment with the variables described in Table 9.
 - *rain_water_isotopes*: contains one csv file per catchment with the variables described in **Table 10**.
- *shapefiles:* contains three subfolders.
- 355 o camels_ch_del: contains two shapefiles. One shapefile includes the derived catchment boundaries associated with each gauge, and the other shapefile marks the location of the gauge stations. Both files are referenced in the Swiss coordinate system LV95 (sometimes also referred to as CH1903+) and were copied from the original CAMELS-CH.
- 360 o *nawa_trend_del*: provides the alternative delineation shapefile for the NAWA
 361 TREND catchments.

362		 nawa_fracht_del: like the anterior, but for NAWA FRACHT catchments.
363	•	gauges_medatada: contains one csv file covering all the metadata associated with
364		each of the 115 gauging stations that will be described in Table 11 .
365	•	stream_water_chemistry: contains two subfolders.
366		o <i>timeseries</i> : contains two nested sub-sub folders. The csv files in both are
367		organized by time series (rows) and attribute variables (columns), and each
368		column represents one of the four water quality variables as described in
369		Table 1. Both nested subfolders contain 86 files.
370		 daily: contains one csv file per catchment at daily resolution.
371		 <i>hourly</i>: contains one csv file per catchment at hourly resolution.
372		 interval_samples: contains three nested sub-subfolders.
373		 nawa_fracht: contains one csv file per catchment covered (24 files).
374		The rows represent the dates, and each column represents one of the
375		water quality variables, as described in Table 2.
376		 nawa_trend: contains one csv file per catchment covered (76 files)
377		and presents a similar structure as the anterior, but now with each
378		column covering one of the variables in Table 3 .
379	•	stream_water_isotopes: contains two subfolders. Each contains one csv file per
380		catchment with any isotope data. The rows represent the dates, and each column
381		represents either deuterium or oxygen-18 data.
382		 <i>isot:</i> contains one csv file per catchment covered (nine files).

- 383

• *ch-irp:* contains one csv file per catchment covered (11 files).

384 Gauge metadata

385 The gauge metadata csv file contains the basic information to allow a proper use of the dataset. Many attributes are a repetition of those provided by CAMELS-CH. Note that the 386 387 coordinate information on northing and easting is always provided in the Swiss reference 388 system LV95, while the gauge_lon and gauge_lat are provided in WGS84. Additionally, due to 389 the potential location difference between the measurement point of the CAMELS-CH 390 streamflow gauge and both NAWA FRACHT and NAWA TREND, further fields were added to 391 ensure consistency when using the data.

392 The attributes gauge_name_{}, gauge_easting_{}, gauge_northing_{} and area_{} refer to 393 specific information from either NAWA FRACHT or NAWA TREND when applicable. The 394 attribute **area_swiss_perc** represents the percentage of the upstream catchment area located 395 in Switzerland and might be useful for users when using the catchment aggregated data.

396 The field **foen_{}_dist** represents the distance in kilometres between the CAMELS-CH 397 streamflow gauge and the NAWA FRACHT or NAWA TREND measurement points (when 398 applicable). Additionally, we also added a correction factor (q_nawat_corrector) for the 399 NAWA TREND measurement points, which can be used to correct the streamflow discharge 400 (as provided in CAMELS-CH) to the new catchment area when using the chemistry data. Finally, 401 the field remarks summarize additional potential information about the gauges that should 402 be considered before using the data.

403	Table 11. Overview of the gauges metadata structure with their respective variables name,
404	description, and units.

Attribute name	Description	Units
gauge_id*	Catchment identifier according to FOEN notation.	-
sensor_id	The same as gauge_id for stations where water chemistry measurements from continuous measurements data are available	-
nawaf_id	Catchment identifier according to NAWA FRACHT notation.	-

Attribute name	Description	Units
nawat_id	Catchment identifier according to NAWA TREND notation.	-
isot_id	Catchment identifier according to ISOT notation.	-
chirp_id	Catchment identifier according to CH-IRP notation.	
gauge_name*	Gauging station name.	-
water_body_name*	Water body name.	-
gauge_easting*	Gauging station easting.	m
gauge_northing*	Gauging station northing.	m
gauge_lon*	Gauging station longitude.	0
gauge_lat*	Gauging station latitude.	0
area*	Catchment area derived using the FOEN outlet.	km²
area_swiss_perc	Percentage of the upstream catchment area located	%
	in Switzerland. A value of 100 means that the	
	catchment is located completely within Swiss	
	borders.	
Q	Information if discharge time series from CAMELS-CH	yes/no
	is available.	
level	Information if water level time series from CAMELS-	yes/no
	CH is available.	
gauge_name_nawaf	Gauging station name according to NAWA FRACHT.	-
gauge_easting_nawaf	Gauging station easting according to NAWA FRACHT.	m
gauge_northing_nawaf	Gauging station northing according to NAWA FRACHT.	m
area_nawaf	Catchment area derived using the NAWA FRACHT	km²
foon nawaf dist	Distance between the gauging station from CAMELS	km
	CH and the NAWA ERACHT sampling location (0 when	NIII
	both are at the same location)	
gauge name nawat	Monitoring site name according to NAWA TREND	_
gauge_name_name	Monitoring site easting according to NAWA TREND.	m
gauge northing nawat	Monitoring site northing according to NAWA TREND.	m
area nawat	Catchment area derived using the NAWA TREND	km ²
	sampling location.	
foen nawat dist	Distance between the gauging station from CAMELS-	km
	CH and the NAWA TREND sampling location (0 when	
	both are at the same location).	
q_nawat_corrector	Weighting factor available to adjust the streamflow	-
	time series to the NAWA TREND catchment area.	
remarks		-

405 * This information is the same as already provided in CAMELS-CH⁵.

406 **Catchment delineations metadata**

The delineated geometry of each catchment is stored in the catchment layer. This layer includes the **gauge_id** field, which is also used for the gauges, allowing for a link between the two datasets. Additionally, the catchment layer also has the fields shown in **Table 12**. These fields ensure consistency between the catchment and gauge datasets, facilitating seamless integration and analysis.

412

413 **Table 12.** Catchment delineations metadata structure with their respective variables name,

414 description, and units.

Attribute name	Description	Unite	
Attribute name	Description		
gauge_id	Catchment identifier according to FOEN notation.		
sensor_id	The same as gauge_id for stations where water	-	
	chemistry measurements from continuous		
	measurements data are available.		
nawaf_id	Catchment identifier according to NAWA FRACHT	-	
	notation ¹⁶ .		
nawat_id	Catchment identifier according to NAWA TREND	-	
	notation ¹⁷ .		
isot_id	Catchment identifier according to ISOT notation ¹⁸ .		
gauge_name	Gauging station name		
water_body	Water body name.	-	
gauge_east	Gauging station easting.		
gauge_nort	Gauging station northing.		
gauge_lon	Gauging station longitude.		
gauge_lat	Gauging station latitude.		
area	Catchment area derived using the FOEN outlet.		
area_swiss	Percentage of the upstream catchment area located in		
	Switzerland. A value of 100 means that the catchment is		
	located completely within Swiss borders.		

415 **Technical Validation**

416 Calibration of the sensor data

The devices used to measure the variables available from sensor data are calibrated twice per year; if there was significant deviation from manual measurements, they are corrected accordingly. Whenter needed, calibration of physico-chemical sensors at NAWA FRACHT^{13,15} stations is performed monthly.

421 Water chemistry measurements first "sanity check"

422 We provided a first assessment of the validity of some of the measured variables. Based on 423 previous literature, we formulated three main hypotheses on the expected variable 424 correlations among themselves. We then tested these hypotheses to determine whether the 425 observed water chemistry measurements are consistent with expectations. Here we 426 computed the correlations using the Spearman correlation coefficient (r_s). Our hypotheses are 427 as follows:

- 428 (i) Stream water EC is broadly negatively correlated with mean discharge, as
 429 demonstrated by previous studies^{31,32}.
- 430 (ii) EC is positively correlated to the measurements of major anions, such as Cl^{-} and NO_{3}^{-} . 431 Conductivity measures the ability of the stream water to conduct electricity, which is 432 directly correlated to the amount of dissolved ions^{33,34}.
- 433 (iii) Stream water temperature and oxygen concentration are negatively correlated.
 434 Increasing temperature decreases the solubility of oxygen in water, moreover, the
 435 increase in water temperature leads to an increase in biological activity, which can
 436 consequently reduce the concentrations of dissolved oxygen in the stream water^{35,36}.
- Therefore, we selected the variables: ec25_lab, Cl, NO3_N and q_mean_sensor from NAWA
 TREND; ec20_lab, Cl, NO3_N and q_mean_sensor from NAWA FRACHT; and temp_sensor, and
- 439 O2C_sensor from the continuous measurements data.

Figure 4 shows the histogram of the distribution of the Spearman correlation coefficients (r_s) computed between electrical conductivity and either mean discharge (a), Cl (b) and NO3_N (c) for NAWA TREND and NAWA FRACHT data. Overall, the correlations between electrical conductivity and mean discharge in **Figure 4a** were largely negative, with values close to r_s of -0.50 for the three data sources. NAWA FRACHT only had one station out of 24 with a positive value, while there were 7 out of 76 for NAWA TREND. These findings are aligned with our hypothesis (i).

447 Moreover, **Figure 4b** shows histograms for the correlation between electrical conductivity and 448 Cl⁻, while **Figure 4c** shows the histograms between conductivity and NO_3^- . Both subplots show 449 that most of the stations exhibit a correlation above 0.50, which supports our expectation 450 from hypothesis (ii).



451
452 Figure 4. Histograms of the Spearman correlation coefficient between (a) EC_{lab} and Q_{mean}, (b)
453 EC_{lab} and Cl⁻, and (c) EC_{lab} and NO₃⁻. The different colors in the subplots represent different data
454 sources, i.e., NAWA TREND in blue and NAWA FRACHT in orange.

455 Figure 5 shows the correlation between temperature and oxygen concentration. Figure 5a 456 shows the histogram of the r_s computed for the daily time series of oxygen and temperature 457 for each of the 16 stations with continuous measurement data for these two variables (Table 458 **1**). All correlations were negative, with only one station with $r_s > -0.70$. Figure 5b shows an 459 example of a daily resolution time series of these two variables for the Mellingen gauge (2018) 460 between 01.10.2019 and 30.09.2020. The figure indicated the expected pattern for the two 461 variables, with oxygen concentrations increasing during the colder months and decreasing 462 with rising temperatures during the summer period. Overall, these results corroborate our 463 hypothesis (iii), which suggested a negative correlation between these two variables.



464

Figure 5. (a) Histograms of the Pearson correlation coefficient between the time series of
oxygen concentration and temperature for all 16 stations covered by the continuous (sensors)
data. (b) Daily time series of oxygen concentration and temperature between 01.10.2019 and
30.09.2020 for gauge Mellingen (2018) used as an example of the typical annual course of the
two variables.

Overall, the rough confirmation of the three hypotheses stated in this section can be used as
broad indication of the reliability of the current water chemistry datasets provided in CAMELSCH-Chem. We acknowledge that this section does not contain a complete validation of the

dataset, yet we believe that it is enough as a first sanity check of the overall validity of CAMELS-CH-Chem.

475 Ionic mass balance for NAWA FRACHT stations

In this section we present the mean, and the 25th, 50th and 75th percentiles of ionic mass 476 balance for all 24 NAWA FRACHT stations, based on the computation of the ionic mass balance 477 478 for each measurement. The full description of the method, along with detailed information 479 about the respective ionic balance of each station, is available in the **Supplementary Material**. 480 The stations presented a maximum 75th values of 4.3 and 70% of them presented a mean ionic balance below 1.5%. Established literature³⁷ suggests that values below 5% ionic 481 imbalance are typically deemed acceptable, while discrepancies exceeding 10% may suggest 482 483 anomalies in measurement or incomplete data. Hence, the present analysis can be used as a 484 validation of most of the NAWA FRACHT measurements. Figure 6 shows the spatial 485 distribution of such values.

487

486

Figure 6. Distribution of the long-term (a) mean, (b) 25th percentile, (c) 50th percentile, and (d)

489 75th percentile ionic mass balance for the NAWA FRACHT stations.

490 **Distance between measurement stations**

491 Regarding the distance between the streamflow measurement gauges and the NAWA FRACHT, 492 16% of the stations (4 out of 24) were more than 5 km away from the respective CAMELS-CH 493 outlet. The maximum distance was 10 km for station 2068, located in the very south of 494 Switzerland (on the Ticino River), with an overall catchment area of 1613.3 km². Furthermore, 495 for NAWA TREND, 14% of the stations (10 out of 72) were more than 5 km away from the 496 CAMELS-CH outlet. Only two gauges had a distance greater than 10 km between the gauge 497 and the sampling location. The station with the maximum distance (20 km) is 2288, located at 498 the Rhine River, and with an overall catchment area of more than 11,000 km².

499

500 These results suggest that the distance between the discharge station and the water chemistry 501 measurement locations of either NAWA FRACHT or NAWA TREND might be neglectable for 502 most basins. For stations where the derived catchment area is considerably different, we 503 suggest users to use the **q_nawat_corrector** to correct the discharge.

504 Isotopes measurements validation

505 The isotopic composition of water samples, specifically δ^2 H and δ^{18} O, was analyzed to assess 506 potential deviations from the Global Meteoric Water Line (GMWL). The GMWL serves as a 507 reference for the isotopic compositions of meteoric water, following **Eq. (1)**. This step was 508 included in the CAMELS-CH-Chem validation phase to demonstrate the usability of the 509 collected data for future users.

δ²H = 8 δ¹⁸O + 10 ‰

510 where δ^{2} H is the deuterium fraction and δ^{18} O the Oxygen-18. Both measurements are in 511 permille (‰) notation according to VSMOW.

Eq. 1

Figure 7 shows the individual subplots **A** to **T** for the total 20 stations with isotope data (nine from ISOT and 11 from CH-IRP), with the δ^2 H and δ^{18} O values plotted alongside the GMWL. The axis limits were set on the basis of the observed range of isotope values across all nine datasets. This comparative approach allowed for a clear identification of any deviations from the GMWL and provided insight into potential fractionation processes in the catchments.

517 Overall, the alignment of the isotope data with the GMWL showed low deviation, with only 518 few samples with apparent evaporative fractionation, indicating that this data provide a 519 reliable basis for future hydrological studies in the provided catchments. For CH-IRP, users can 520 also refer to their original publication, where Staudinger, 2019¹⁹ also provides lab standards 521 and errors for their measurement stations, along with potentially problematic measurements.



523 Figure 7. Dual isotope plots of δ^2 H and δ^{18} O for the nine ISOT and 11 CH-IRP sampling locations

524 in reference to the Global Meteoric Water Line (GMWL, dashed orange line). Blue circles (dark

525 for ISOT and light for CH-IRP) represent individual water samples covering their respective 526 entire timeseries.

527 Code Availability

528 The code used to produce the current dataset is available at: <u>https://github.com/camels-</u> 529 <u>ch/camels-ch-chem</u>. The scripts are organized to enable users to follow a logical sequence 530 during code usage. Finally, the code used to derive all figures and the technical validation is 531 available at <u>https://github.com/thiagovmdon/camels-ch-chem_paper</u>.

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537 Author contributions

538

539 The co-authors T.N. and M.H. initiated and coordinated the project. U.S., M.H. and T.N. 540 collected and pre-processed the stream water chemistry, isotopes from ISOT and atmospheric 541 deposition data. S.P. collected the atmospheric deposition data. R.S. collected and processed 542 the agricultural data. M.K., collected and processed the land cover and the rainwater isotopes 543 data. M.S. collected and processed the isotopes data from CH-IRP. F.S. and P.R. offered 544 guidance regarding the FOEN data. T.N. and M.K. wrote the data aggregation and processing 545 codes in Python. R.S. and U.S. processed the catchment boundaries. M.K., P.H., M.F. and J.S. 546 contributed with some dataset references and discussion. T.N and R.S. made all figures. T.N. 547 wrote the first draft. F.F. retrieved the funding for the project. All co-authors participated in 548 reviewing the manuscript.

549 **Competing interests**

550 The authors declare no competing interests.

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646 Supplementary material

647 Continuous measurements data

Jonunuous	measu	emento	uald

Name of	Years	Manufacturer/Type	Accuracy	Remarks
attribute			according to	
			manufacturer	
temp_sens	1976 -1991	Hartmannn u. Braun / CMR	0.2°C	-
or		TEUC with PT100 (3		
		conductors)		
		Camille Bauer / Eurax V601	0.15 + 0.02	-
	since 1991	with PT100 (3 Leiter)	°C/°C	
		Camille Bauer / Eurax V604	0.15 + 0.02	-
	since 1991	mit PT100 (3 conductors)	°C/°C	
		Rotax PT100 3-conductors	0.15 + 0.02	-
	since 2002	resistance thermometer	°C/°C	
		Rotax PT100 3-conductors	0.15 + 0.02	-
	since 2011	resistance thermometer	°C/°C	
	since	Rotax PT100 4-conductors	0.1°C +0.0017	-
	2018/19	resistance thermometer	°C/°C	
ec_sensor	1976 - 1991	Wösthoff / EMHD 1	5 - 10 μS/cm	-
	since 1991	Siemens / SIPAN 4EL	4 - 7 μS/cm	-
	since 2002	Quadroline LF296 with	0.5% + 1Digit	-
		Tetracon 700		
	since	Endress+Hauser / Memosens	< 4 %	-
	2018/19	CLS82D		
	since 2022	WTW TetraCon 700 IQ	< 4 %	only
				Andelfinge
				n

				hydro_id_2 044, Rekingen hydro_id_2 143, Weil hydro_id_2 613
O2C_sensor	1976 - 1981	WTW / OX1 39	0.23 mg/l	-
	1981 - 1991	Orbisphère / Modèle 2116	0.2 mg/l	-
	2005 /2006	Hach-Lange / LDO oxygen measurement	± 0.2 mg/l	-
	since 2018/19	Endress+Hauser / Oxymax COS61D	± 2 %	-
pH_sensor	1976 - 1991	Hartmann u. Braun / UPY3	0.02	-
	since 1990	Jenco / Modell 6300N	0.10%	-
	since 1993	Endress+Hauser / Mycom CPM 121 mit Ceratex CPS 31	0.03 - 0.04	-
	since 2018/19	Endress+Hauser / Memosens CPS31D	< 0.05	-

649

650 **Supplementary Table 1.** Overview of the manufacturer type, accuracy and geenral remarks

651 for the intruments usef for obtain the continuous measurements data.

652 NAWA FRACHT (previously called NADUF)

- 653 Details are in a separate table depending on station, year, and parameter.
- 654

Name of attribute	Units	Method	min loq	max loq
alk	mmol/l	acidimetric titration (automated)	0.02	0.2
As	μg/l	ICP-MS	0.5	0.5
Ва	µg/l	ICP-MS	0.5	0.5
Br	mg/l	IC	0.01	0.05
Cd	μg/l	ET AAS / ICP-MS	0.01	0.02
Ca	mg/l	calculated from hardness minus magnesium		
Cl	mg/l	CFA (photometric) / IC	0.5	1.5
Cr	µg/l	ET AAS / ICP-MS	0.1	0.5
Cu	μg/l	ET AAS / ICP-MS	0.05	0.5
doc	mg/l	combustion-infrared / wet- oxidation-infrared	0.1	0.5
drp	mg/l	CFA (photometric) / photometric	0.001	0.005
ec25_online	μS/cm			
ec20_lab	μS/cm			
F	mg/l	IC	0.01	0.05
Fe	mg/l	ET AAS / ICP-MS	0.1	0.5
Pb	μg/l	ET AAS / ICP-MS	0.1	0.2
Mg	mg/l	AAS / ICP-AES / IC	0.05	1
q_online	m³/s			
Hg	μg/l	Cold-vapour-AAS	0.002	0.05

Ni	µg/l	ET AAS / ICP-MS	0.1	0.5
NO3_N	mg/l	CFA (photometric) / IC / photometric	0.005	0.2
O2C_online	mg/l			
O2S_online	%	calculated from oxygen concentration, water temperature and elevation		
pH_lab	-			
pH_online	-			
к	mg/l	AAS / ICP-AES / IC / IC	0.05	0.5
H4SiO4	mg/l	CFA (photometric)	0.35	1
Na	mg/l	AAS / ICP-AES / IC / IC	0.05	2
Sr	μg/l	ICP-MS	5	5
SO4	mg/l	titration / FIA (photometric) / IC	1	5
tfp	mg/l	persulfate digestion	0.003	0.003
th	mmol/	EDTA titration (automated)	0.09	0.2
tn	mg/l	persulfate digestion / combustion- chemiluminescence	0.1	0.5
toc	mg/l	combustion-infrared	0.1	0.5
tp	mg/l	hydrogen peroxide digestion / Persulfate digestion	0.003	0.02
tss	mg/l	filtration and weighing	1	1
temp_online	°C			
Zn	μg/l	AAS / ICP-MS	0.1	1

655 **Supplementary Table 2.** Overview of the methods and intruments accuracy used obtaining 656 the NAWA FRACHT data.

657 NAWA TREND

These data are measured by cantonal authorities according to the Swiss Modular Stepwise Procedure (https://modul-stufen-konzept.ch/en/nutrients/). No detailed measurement methods are recorded, only the range of the limits of quantifications. Detailed limits of quantifications depending on stations are given in the appendix.

Name of	Units	min loq	max loq
attribute			
NH4_N	mg/l	0.002	0.096
Cl	mg/l	0.005	5.6
q_canton_max	m³/s		
q_canton_mean	m³/s		
q_canton_min	m³/s		
doc	mg/l	0.1	1
drp	mg/l	0.001	0.05
ec25_lab	μS/cm	0.3	10
ec25_online	μS/cm		
q_online	m³/s		
NO3_N	mg/l	0.002	1
NO2_N	mg/l	0.0006	1
O2_lab	mg/l	0.1	0.5

O2_online	mg/l		
O2S	%		
pH_lab	-	1	1
pH_online	-		
temp_lab	°C		
temp_online	°C		
turbidity_online	NTU		
tn	mg/l	0.041	1
tp	mg/l	0.0008	1.7

662 **Supplementary Table 3.** Overview of the intruments accuracy used obtaining the NAWA

663 TREND data.

664 Ionic balance of NAWA FRACHT data

Gauge id	Ionic mass balance (%)					
	Mean	P25	Median	P75		
2009	0.435985	-0.545902	0.359950	1.299944		
2016	1.639680	0.733901	1.594390	2.505173		
2018	1.205843	0.271829	1.278908	2.133413		
2044	2.099322	1.391718	2.050711	2.778935		
2067	0.398352	-1.059095	0.216745	1.551320		
2068	1.320987	0.111855	1.115087	2.224546		
2085	1.191098	0.441571	1.173093	1.783882		
2106	2.179472	1.479517	2.168280	2.797340		
2112	0.522289	-0.073699	0.377360	1.071219		
2130	1.289813	0.740609	1.562212	2.164777		
2135	1.844861	1.389279	1.911113	2.438441		
2143	1.321940	0.553701	1.265323	2.021457		
2174	0.628014	-0.233800	0.510192	1.335572		
2179	1.292728	0.817671	1.239803	1.714320		
2243	1.544782	0.565584	1.515227	2.398968		
2386	2.531875	2.043512	2.423313	3.009654		
2415	3.148087	2.032618	2.854008	4.268473		
2462	-0.064403	-0.901463	-0.148776	0.720982		
2467	1.449764	0.803557	1.405023	2.055323		
2473	0.587801	-0.299268	0.457965	1.363267		
2608	1.311412	0.905194	1.240102	1.643632		
2613	1.329235	0.510777	1.241231	2.178189		

665 **Supplementary Table 4.** Mass balance statistics for each of the stations with NAWA FRACHT 666 measurements.

667 Additional stations

668 NAWA FRACHT: Alpthal stations <u>https://opendata.eawag.ch/dataset/naduf-national-long-</u> 669 <u>term-surveillance-of-swiss-rivers-2024-2</u>