# *Note: All line number references in author responses refer to the track-changes document, unless otherwise indicated*

#### **Reviewer 1**

Overview

This manuscript presents a study of thermokarst lakes in the Northern Territories, Canada, investigating hydrological, meteorological, and watershed controls on the lakes' water balance. Twenty-five lakes were sampled five times over the spring-summer season in 2018 and analyzed for stable oxygen and hydrogen isotope ratios. The isotope data were used to calculate the evaporation-to-inflow (E/I) ratio and the isotopic composition of lake source water ( $\delta$ I). Four water balance phases were identified, associated with shifts in meteorological and/or hydrological changes. The E/I and  $\delta$ I were compared to meteorological and watershed data (e.g., precipitation amount and watershed area to lake area ratio (WA/LA)). Subsequently, the relationship between WA/LA and E/I was used to estimate the average E/I of 7340 lakes in this region, finding that the lakes are not vulnerable to desiccation, and that lakes with smaller WA/LA are likely to be more influenced by increased evaporation in a future warmer climate.

General comments

The manuscript presents an interesting approach to evaluate water balance changes and to estimate E/I, which is relevant for the scope of HESS. The manuscript is well written with an easy-to-follow structure and clear figures. However, there are some issues with the dataset size and associated assumptions that need to be more clearly outlined. I also have some questions concerning the chosen method to infer  $\delta I$ . I have listed my concerns and suggestions for improvement/clarifications below.

• Thank you for your thorough review of our manuscript. We believe that we have addressed all your comments and have implemented most of the suggestions you made.

Precipitation dataset size: I have some concerns about the small size of the precipitation dataset used for the "isotopic framework". Are 11 snow samples and 13 rain samples enough to produce a reliable LMWL and estimate  $\delta P$ , or would it be better to use a more conservative approach and use the GMWL? How well do the precipitation samples cover the annual range of precipitation isotopic variability in this region? And why not compare your data to GNIP data from Inuvik (from the 1980s)? Justify your approach to use local precipitation data (because they are most representative for the study area and cover the same period as your lake water dataset?). How does the uncertainty related to the precipitation data affect the uncertainty in the  $\delta^*$  and  $\delta I$  calculations? Some more information about the precipitation samples would be useful to clarify how well estimated  $\delta S$ ,  $\delta R$  and  $\delta P$  are. Were the snow samples collected soon after snowfall, or can the "end of winter snowpack" have experienced post-depositional fractionation processes (e.g., sublimation) before sampling? Were the liquid samples event-based and/or representing all precipitation events from May to September 2018? Have the precipitation isotope data been amount-weighted? Please clarify.

- We have clarified the collection of snow samples. They were not collected during the winter as precipitation fell, but the end of winter snowpack was sampled by pushing a cylinder vertically through the snowpack at the end of winter, putting the sampled snow into a plastic bag to melt and then transferring the melted snow into a sampling bottle (L107-108). In this way, we incorporate any fractionation processes that may have occurred as the snow was on the ground and measure the amount-weighted snow isotope composition by virtue of how we sampled the snowpack.
- We relied on the precipitation samples we collected to establish the LMWL because they most closely represent the input water to the lakes (compared to the GMWL), and we believe we have sufficiently sampled both rainfall and snowfall to determine the LMWL, at least for the year of 2018. Below we explain why we will remain with the LMWL we developed:
  - Unfortunately, there are only a few years of GNIP data available from Inuvik, from 1986-1989, with many months only having one year of data, and this is so long ago that we worry the values for  $\delta_S$  and  $\delta_R$  may have changed since the eighties as climate change alters global atmospheric circulation and reduces sea ice in the Beaufort Sea. Although, the equation of the GNIP LMWL line and our LMWL line are similar:
    - GNIP LMWL equation  $\delta^2 H = 7.3 \times \delta^{18} O 3.5\%$ ,
    - Our LMWL  $\delta^2 H = = 7.1 * \delta^{18} O 10.0\%$
  - $\circ \quad By \ complete \ coincidence, \ another \ group \ has \ developed \ a \ LMWL \ for \ Inuvik \ using \ three \ years \ of \ data \ and \ 134 \ precipitation \ samples \ between \ 2015 \ and \ 2018 \ (Fritz \ et \ al., \ 2022). \ The \ LMWL, \ \delta_R \ and \ \delta_S \ values \ they \ calculate \ are \ very \ similar \ to \ ours:$ 
    - Fritz et al. (2022) values for  $\delta_{R}$ ,  $\delta_{S}$ , LMWL
      - Average, non-amount weighted  $\delta_R$ : -16.9‰ ( $\delta^{18}$ O), -131.8‰ ( $\delta^{2}$ H)
      - Average, non-amount weighted  $\delta_s$ : -23.8‰ ( $\delta^{18}$ O), -181.7‰ ( $\delta^{2}$ H) (Fritz et al., 2021).
      - LMWL:  $\delta^2 H = 7.4 * \delta^{18} O 6.7\%$
    - Our values
      - Average, non-amount weighted  $\delta_R$ : -17.03‰ ( $\delta^{18}$ O), -129.54‰ ( $\delta^{2}$ H)
      - Average, non-amount weighted  $\delta_s$ : -24.61‰ ( $\delta^{18}$ O), -184.19‰ ( $\delta^{2}$ H)
      - LMWL:  $\delta^2 H = 7.1 * \delta^{18} O 10.0\%$
  - $\circ$  Given the close agreement in the LMWL,  $\delta_R$  and  $\delta_s$  values between our values and Fritz et al. (2022), we believe that redoing the analysis using the Fritz. et al (2022) data would not cause a meaningful difference in the results. We now discuss in the manuscript that our LMWL fits closely with the Fritz et al. (2022) and GNIP LMWL (L136 143).
- Post-depositional fractionation of snowfall was likely minimal (approx. 1‰ δ<sup>18</sup>O) See Ala-aho et al. (2021), where they compare snowfall samples to corresponding snow layers at the end of winter for an Arctic tundra site (Figure 9a, 10a). Again, our agreement with Fritz et al. (2022) also suggests there was little change in snow isotope composition.
- In regard to precipitation data affecting the uncertainty in  $\delta^*$  and  $\delta_I$  calculations:  $\delta^*$  could impact  $\delta_I$  through precipitation samples since we assume that atmospheric vapour ( $\delta_{As}$ ) is in equilibrium with the isotope composition of summertime precipitation (Equation A5). Since we seem to capture  $\delta_R$  well when compared to Fritz et al. (2022), we think that we have likely calculated  $\delta_I$

well. Also, lakes seem to plot around the LEL, which we calculate independently of lake water samples, suggesting that we have estimated  $\delta^*$  accurately.

• We have clarified the calculation of  $\delta_s$  and  $\delta_R$  and that they are the average of all samples without any weighting by precipitation amount (L134 – 135).

Analytical uncertainty: Are duplicate measurements of every fifth sample enough to determine the analytical uncertainty? In many labs, each sample is injected multiple times (sometimes more than 10 times), the first replicate(s) discarded, and the rest used to calculate an average. How did you deal with drift and memory effects?

- By duplicate measurements, we do not mean duplicate injections. Each measurement consists of 8 injections of ~1000nL, of which the first two were discarded, with the remaining six injections averaged for the measurement.
- VSMOW and VSLAP standards provided by the instrument manufacturer were interspersed throughout the sample run to confirm the accuracy of the instrument.
- We have added this detail to the methods section (L117 125).

Isotope framework: I generally like the approach to present an "isotope framework", but some clarifications are needed when comes to presentation of isotope data and the used terminology. One example is the inconsistent use of the delta notation ( $\delta$ ). Some examples of this are on line 44 where  $\delta$  is missing when introducing water isotope analysis (which is the analysis of the stable isotope ratios  $\delta$ 180 and  $\delta$ 2H), and ' $\delta$ - $\delta$  space' which should be ' $\delta$ 2H- $\delta$ 180 space'. There is also a parenthesis missing in equation 1, which should be written:  $\delta$ sample = (Rsample/RVSMOW-1) \* 103. These fundamental things need to be stated correctly. Furthermore, it is not clear when  $\delta$ I is referring to  $\delta$ 180I and when/if it refers to  $\delta$ 2HI as well?

- We have corrected it to " $\delta^2$ H- $\delta^{18}$ O space" throughout the ms.
- We now clarify that  $\delta_I$  refers to both isotopes, and  $\delta^{18}O_I$  refers to just  $\delta^{18}O$  (L155). We use only  $\delta^{18}O_I$  values in the results and discussions since  $\delta^{18}O_I$  perfectly correlates with  $\delta^2H_I$  because  $\delta_I$  values are bound to the LMWL.
- We have corrected the  $\delta_{\text{Sample}}$  formula (L116), however we have removed the "\*10<sup>3</sup>" since a reviewer in another manuscript has pointed out to us that  $\delta$  notation is not inherently in ‰ units (Coplen, 2011). This and the other manuscript will be a part of a PhD thesis, so we would like to keep the formulas consistent between the chapters.

Approach to calculate  $\delta I$ : Have you considered using the more recent 'MWL source implementation' method by Bowen et al. (2018) to model  $\delta I$  values? Using this approach, you supply a MWL equation (you could test the GMWL and your LMWL to see how much difference it makes) and a hypothesized LEL slope with confidence intervals, as well as your lake water  $\delta 2H$  and  $\delta 18O$  values with uncertainties.

• We were aware of this approach and but not considered it. Our understanding of the benefits of the MWL source implementation approach is that it avoids biasing the  $\delta_I$  of lakes when a regression through lake water samples is used to calculate the LEL, while there is correlation between evaporation-to-inflow ratios and  $\delta_I$ . Since we calculate  $\delta_I$  using a fixed point in  $\delta^2$ H- $\delta^{18}$ O space that is calculated independently of lake water samples, and calculate a lake-specific evaporation line for each lake water sample, we believe that our method is not vulnerable to the errors that the Bowen et al. (2018) approach is meant to address. Our understanding is that the

approach introduced by Bowen et al. (2018) is more geared towards large-scale studies covering larger areas than our region, and they actually encourage "model-based approaches" for site-specific studies like ours "*We also believe that* [a model-based] *approach should be more frequently adopted in site-based studies as an alternative to regression*..." (p1028, Bowen et al., 2018). Therefore, we will remain with our initial approach to calculating  $\delta_{I}$ , but we will keep this other method in mind for our future research.

Specific comments

Lines 18-19: Rephrase the sentence starting "WA/LA strongly predicted average lake E/I ratio (R2 = 0.74)..." to "Lakes with smaller WA/LA tended to have higher E/I ratios (R2 = 0.74) because they received relatively less inflow. I think "strongly predicted" is an exaggeration.

• We disagree that it is an exaggeration – a single variable explaining three quarters of the variability in a complex environmental system is uncommon. However, we changed the wording anyways.

Line 40: evaporation-to-inflow ratios. Use this wording throughout the paper (sometimes it says evaporation to inflow, sometimes evaporation/inflow ratio).

• We have changed the wording throughout the manuscript to be consistent.

Line 44: Change to "In several studies, stable water isotope ( $\delta$ 18O and  $\delta$ 2H) analysis..." and add a couple of references.

Changed to: "In several studies, water isotope (δ18O and δ2H) analysis has been the primary method used to efficiently characterize the water balance of a large number of thermokarst lakes because the isotope composition provides an integrated measure of influential hydrological processes (Gibson, 2002; Edwards et al., 2004; Turner et al., 2014; Narancic et al., 2017)." (L39)

Line 60: Explain the concept "snowmelt bypass" here. Now it isn't explained until in the methods (lines 88-89).

• Added: "Snowmelt bypass occurs when less dense (~0°C) freshet runoff flows underneath lake ice and passes through a lake without mixing with and replacing the deeper and denser (<4°C) lake water." (L63 – 64)

Line 74: Remove "All lakes we selected were either headwater lakes or downstream of lakes that we sampled", since this information is repeated on lines 76-77.

• Change implemented. (L79 -80)

Line 87: How much did the first day of ice-free season vary between the lakes?

Added: "June 15 was chosen as it marked the first day of the ice-free season for most lakes and was intended to capture the influence of snowmelt bypass, however the southernmost lakes (Lakes 7-12) became ice-free on June 7, and some of the northernmost lakes (Lakes 49-55) became ice-free on June 17 or June 18." (L96 – 98).

Lines 88-89: Move the definition of snowmelt bypass to the introduction and rephrase it. This sentence does not read well and needs to be clarified.

• Change implemented. (L94-95)

Line 97: Explain "end-of-winter snow". Does this mean that you sampled the snow in the end of winter soon after it fell, or that you collected a core of snow accumulating over a longer period? If the latter, how do post-depositional processes impact the isotope values?

- There is likely minimal change in snow isotope compositions after snow is deposited, as we describe above. See Ala-aho et al. (2021).
- Also as we describe above, we have clarified that the snow samples are taken using a vertical core through the end-of-winter snowpack. (L107-108).

Line 98: Are  $\delta S$  and  $\delta R$  amount-weighted? Change to  $\delta S$ now and  $\delta Rain$  throughout the text to match the terms used in Figure 3 (or change to  $\delta S$  and  $\delta R$  in the figure).

• They are not amount weighted; we have made the terms similar between the text and figures.

Line 107: Equation misses parentheses around (Rsample/RVSMOW - 1)

• As described above, we have changed the formula to omit the  $x10^3$  now as  $\delta$  notation is not inherently in ‰ units. (L116).

Lines 113-114: When introducing the "fundamental linear relationships", describe the global relationship (i.e., the GMWL) as well. This could also be added to Figure 3.

- We now describe the GMWL and have added it to Figure 3.
- Added: "The isotope composition of precipitation from anywhere in the world tends to plot closely to the Global Meteoric Water Line, a strong linear relation in  $\delta^{18}O$ - $\delta^2H$  space which is defined as  $\delta^2H = 8.0 * \delta^{18}O + 10$  (Craig, 1961)." (L129 131).

Line 114: Change 'δ-δ space' to 'δ2H-δ18O space' (also on line 122).

• Change implemented throughout the ms.

Line 116: Is  $\delta P$  the average between the two  $\delta S$  and  $\delta R$  values, or the average of the full range of rain and snow values?

- It is the average of the two ( $\delta_s$  and  $\delta_R$ ) values, otherwise the number of  $\delta_s$  and  $\delta_R$  samples would weight  $\delta_P$ . We have clarified this now in the text.
- Added: "The average isotope composition of rainfall ( $\delta_R$ ) and snowfall ( $\delta_S$ ) were calculated by averaging the isotope composition of all the rainfall and snowfall samples we collected respectively, with  $\delta_P$  representing the average of  $\delta_S$  and  $\delta_R$ ." (L134 135)

Lines 116-122: As it is explained now, it is not clear how the LEL was defined. Looking at Figure 3, the slope of the LEL is different between  $\delta P$  and  $\delta SSL$  compared to between  $\delta SSL$  and  $\delta^*$ ? It also says that  $\delta SSL$  is located along the LEL. Which line is the LEL equation referring to? Why is it not a straight line from  $\delta P$  to  $\delta^*$ ? Please clarify.

- We double checked our calculations and found an error: we had calculated  $\delta_{SSL}$  for a lake fed with  $\delta_R$  instead of  $\delta_P$ , which caused the bend in the line. We didn't initially notice this error because there is usually a slight bend in the line associated with uncertainties in the fractionation equations for  $\delta^2 H$ , which influences  $\delta_{SSL}$  and  $\delta^*$  estimates. The line has now been fixed by calculating  $\delta_{SSL}$  using  $\delta_P$  as the source water. (Figure 3, p10)
- We have now labelled the line referencing the LEL on Figure 3. (p10)
- Modified the description of the LEL for clarity: "Evaporated waters tend to plot along a Local Evaporation Line (LEL), which can be defined independent of measured lake water isotope compositions. Using this approach and for the case of a lake fed by waters of mean annual isotope composition of precipitation, the LEL was anchored at δ<sub>P</sub> and the theoretical isotope composition of a lake at the point of total desiccation (δ<sup>\*</sup>), which is dependent on air temperature, relative humidity and the isotope composition of atmospheric moisture in the region (Gonfiantini, 1986). Along the LEL exists a useful reference point where the amount of evaporation a water body is experiencing is equal to the amount of water input, defined as δ<sub>SSL</sub> (steady-state lake water isotope composition of a terminal basin)." (L144 150)

Line 122: What does SSL stand for? Add reference for  $\delta$ SSL definition.

• Steady state lake, we have added this now: "defined as  $\delta_{SSL}$  (steady-state lake water isotope composition of a terminal basin)." (L149 – 150)

Lines 126-128: Describe the  $\delta$ I calculation here as well (not only in the appendix).

• We now split the paragraphs describing the calculation of  $\delta_I$  and E/I, and more clearly describe how  $\delta_I$  is calculated. (L144 – 166)

Line 131: "where  $\delta L$  is the isotope composition of the lake water and  $\delta E$  is the isotope composition of evaporated vapour from the lake (Gonfiantini, 1986)"

• Added. (L165)

Line 136: cumulated rainfall?

• Total rainfall amounts within the periods of time being discussed.

Lines 150-151: What do you mean by "data were transformed if the distribution was non-uniform"?

- If the distribution of the data on a histogram was non-uniform, a mathematical function (e.g. ln) was applied to the data in order to transform the data into a uniform distribution.
- Changed to: "The distribution of each variable was plotted on a histogram and data were mathematically transformed if the distribution was non-uniform." (L184 185)

Line 152: Rephrase this sentence (removing the "strong") and add the R2-value.

• Change implemented. (L187)

Line 170: This is the first time you use  $\delta$ 180I. Does  $\delta$ I mentioned throughout the paper refer to  $\delta$ 2HI and  $\delta$ 180I collectively, or  $\delta$ 180I? Please clarify.

- We now clarify this when we introduce  $\delta_I$  in the methods section
- Changed to: "Isotope compositions of the lakes were used to estimate the average isotope composition of source water ( $\delta_I$ , or  $\delta^{18}O_I$  when referring to just  $\delta^{18}O$ )" (L155)

Line 170: "We observed distinct shifts in lake water isotope composition along the LEL..." – This is not easy to read from Figure 3. Do you mean for individual lakes, or the lakes in general? It could be interesting to indicate which lake is which, to be able to compare the different lakes' responses during P1-P4. This could for example be done (in a supplementary figure?) by giving the lakes different colors, and assign each sampling date a different symbol, and/or by drawing lines between data points from the same lake.

- We explored some of the lake-specific changes in water isotope composition in our data analysis, but it becomes very messy to plot all lakes at once, and most lakes followed the general pattern over time that we describe in the discussion (the four water balance phases). There are some interesting patterns for a couple lakes and for clusters of lakes where we have some upstream and downstream water samples, but overall, our goal with this paper was to generally describe the variability in lake water balances over space and time. Maybe in a future manuscript we will explore more lake-specific water balance evolutions, and the impact of upstream lakes (and their watershed properties) on downstream lakes.
- Changed to: "In general, we observed distinct shifts in lake water isotope composition..." (L206)

Lines 174 and 180: Remove "very".

• Change implemented. (L210, 216)

Lines 178 and 184: Change "compositions" to "values"

• Change implemented. (L214, 220)

Lines 193-194: "On September 3, some lakes plotted close to the LMWL, indicating that their waters had experienced negligible amounts of evaporation (Figure 3)." Does this mean that the same lakes plotted on the LEL before, and that the high precipitation amounts during P4 "reset" the lake water to be closer to  $\delta I$  by removing the old evaporation signal?

• Yes, their waters had no evaporation signal. These lakes likely had large WA/LA ratios and short water residence times as a result, making their fluctuations in E/I more sensitive to short-term changes in meteorological conditions.

Line 196: Two thirds of the lakes.

• Change implemented. (L232)

Line 221: Remove "results" before "appears".

• Change implemented. (L259)

Lines 245 and 246: Change "cooler" to "lower".

• Change implemented. (L283, 284)

Line 278: Earlier you mention only 5 downstream lakes (e.g., in Table 1). Where did the 6th lake come from (also presented in Figure 5)?

• We had incorrectly labelled Lake 12 in our table as a headwater lake when it is in fact a downstream lake. We have corrected the table and text accordingly.

Line 320: evaporation

• Change implemented. (L366)

Line 321: led

• Change implemented. (L367)

Line 329: remove "as high as"

• Change implemented. (L375)

Line 357: Is  $\delta Ps$  the value referred to as  $\delta P$  in Table A1, or where do you present that value?

•  $\delta_{Ps}$  is summertime precipitation, while  $\delta_P$  is all precipitation. This is clearer now in our revised Table A1, where we describe all the parameters in the isotope framework and how they are generated.

Figure 1: Add overview map (e.g., Canada), and add black triangles and red square to legend.

• Change implemented.

Figure 2: Add reference to meteorological data. Clarify that you show the cumulative precipitation amounts. The sample days are shown by vertical dashed, not dotted lines. Mean daily air temperatures are indicated by horizontal dashed lines.

• Changes implemented.

Figure 3: Please explain all elements in the caption, e.g., that the  $\delta$ Snow,  $\delta$ Rain (which are called  $\delta$ S and  $\delta$ R in the text) and  $\delta$ P are averages (it looks like  $\delta$ Rain displays the median, but maybe the average and median are very close?), what the box and whisker plots show, and that the numbers refer to sampling dates. It would also be good to add the GMWL for reference. Which line does the LEL equation refer to? The line between  $\delta$ P and  $\delta$ SSL and the line between  $\delta$ SSL and  $\delta^*$  have different slopes. Why? Change color for the snow samples or the May 1 samples, as it is confusing that both are displayed in grey. You could also use different symbols for precipitation samples and lake samples, to make them easier to differentiate.

- We have added descriptions for all elements shown in Figure 3. The colour of snow samples has been changed to purple, and the symbols for precipitation have been made different to the symbols for lakes. A legend has also been added for all the different isotope samples.
- As we mentioned above, we had made an error in calculating  $\delta_{SSL}$ . The LEL is now a straight line. The calculation of  $\delta_{SSL}$  is now clarified in Table A1.

Figure 4: It says four sampling dates in the caption but should be five. What do you mean by "lake-specific" change in E/I in (c)? Why not also present a panel with the measured  $\delta$ 18O values, to see how much these values differ from  $\delta$ 18OI?

- We have corrected the number of sampling dates.
- We have clarified what "lake-specific change in E/I" means, we calculated the difference in E/I between sampling points for each lake, and then show the distribution of those values in (c). We added the equation for this to the figure caption.
- We are not sure of the reason to compare  $\delta^{18}$ O to  $\delta^{18}$ O<sub>I</sub>, we already show the distribution of  $\delta^{18}$ O values for each isotope sample type in Figure 3, and one can get an idea of  $\delta_I$  values by observing how lakes plot above or below the LEL.

Figure 6: Are the temperatures and precipitation values in from the year of lake water sampling or the 1980-2020 values? And how were the curves in the three lowermost panels generated? The E/I values at the sampling dates are not the same as presented in Figure 4b? And how were the values between the sampling dates interpolated? Please clarify.

• We have now clarified in the figure caption that this is a conceptual diagram and not pure data, although the temperature and precipitation data do represent 1980-2020 averages. We have redrawn the E/I and  $\delta_I$  lines to better represent our data.

Table 1: Change Polygon Extent to Ice-wedge Polygon Coverage.

• Change implemented.

Table 2: Clarify in caption that you mean  $\delta$ 180I. What do you mean by "adjusted R2"?

• Adjusted R<sup>2</sup> is used in cases with multiple explanatory variables, in an attempt to control for the natural tendency for R<sup>2</sup> to increase as additional variables are added. In R, the Wherry formula is used to calculate adjusted R<sup>2</sup>: R<sup>2</sup><sub>adj</sub> =  $1 - (1 - R^2)*((n-1)/(n-p-1))$ , where n is the sample size and p is the number of explanatory variables (i.e., predictors). When there is only one explanatory variable, R<sup>2</sup>  $\approx$  R<sup>2</sup><sub>adj</sub>. We have added this to the caption.

# Reference

Bowen, G.J., Putman, A., Brooks, J.R., Bowling, D.R., Oerter, E.J., Good, S.P., 2018. Inferring the source of evaporated waters using stable H and O isotopes. Oecologia 187 (4), 1025–1039. https://doi.org/10.1007/s00442-018-4192-5.

- References
  - Ala-aho, P., Welker, J. M., Bailey, H., Højlund Pedersen, S., Kopec, B., Klein, E., Mellat, M., Mustonen, K., Noor, K., and Marttila, H.: Arctic Snow Isotope Hydrology: A Comparative Snow-Water Vapor Study, 12, 150, https://doi.org/10.3390/atmos12020150, 2021.

- Coplen, T. B.: Guidelines and recommended terms for expression of stable-isotope-ratio and gas-ratio measurement results, 25, 2538–2560, https://doi.org/10.1002/rcm.5129, 2011.
- Fritz, Michael; Wetterich, Sebastian; McAlister, Joel; Meyer, Hanno (2021): Stable oxygen (δ<sup>18</sup>O) and hydrogen (δD) isotopes in precipitation from Inuvik in the western Canadian Arctic (2015-2018). PANGAEA, https://doi.org/10.1594/PANGAEA.935027 [data set]
- Fritz, M., Wetterich, S., McAlister, J., and Meyer, H.: A new local meteoric water line for Inuvik (NT, Canada), 14, 57–63, https://doi.org/10.5194/essd-14-57-2022, 2022.

# **Reviewer 2**

The manuscript employs the well known isotope mass balance formulations to compute the water balance, namely, the evaporative water loss to total inflow (E/I) ratio in 25 Thermokarst lakes in Canada. The authors subsequently compare the E/I to the lake and catchment geometry and cover [Catchment Area to Lake area ratio, Lake Depth, Vegetation, gradient, Lake area etc]. Interestingly, the E/I ratio of the 25 lakes correlate well (R2=0.74) with the Catchment/Lake area ratio, allowing the authors to compute the E/I ratio for 7340 Thermokarst lakes without needing the hydrologic or isotopic measurement in these lakes.

The 25 lakes have similar range in depth (0-4 m) dictating the volume of the water in the lakes is proportional to their area (because of similarity in lake depth). This would mean the underlying mechanism for the good correlation between E/I and Catchment/Lake area ratio is that the later is a good surrogate for Catchment/Lake Volume and thus to residence time (Volume/Inflow). The major question now is what would happen if the lakes (25 lakes) vertical dimension was significantly variable and if the 7340 lakes have variable depth properties? The validity of the assumption that the lakes have similar depth property must be clearly reflected in the manuscript or the uncertainties of using the approach or extending the approach beyond these lakes needs to be highlighted in the manuscript.

• Thank you for your comments, variability in lake depth, and therefore volume, is not something we had considered. We have done some simple statistics using lake depth data published in a previous study of 34 different lakes in this region by Pienitz et al. (1997). The median lake depth was 3.0 m, average of 5.0 m, and a range between 1.5 to 18.5 m. We now discuss that our E/I estimates may underestimate E/I for these deeper lakes, as increasing lake depth likely increases residence time and therefore the opportunity for lake water to evaporate. It would be interesting for future work to compare these deeper lakes to shallower lakes, with respect to E/I. (L328 – 334)

Other detailed comments (13 comments) are provide in the attached PDF.

#### PDF COMMENTS:

i suggest you delete this

• change implemented (L14)

Did you account for interconnection between the sampled lakes. Given that the upstream lakes feed downstream lakes isotope enrichment in the downstream lake is not only the function of water balance but also the influence of enriched upstream inflow. If you have accounted for this please indicate it in the sentence next to this one.

• We did not account for it in the isotope framework directly, but we do separate headwater and downstream lakes when we present results in Figure 5, and we also only use upstream lakes to upscale Average E/I ratios in Figure 7.

please use consistent notation. In figure 3 for instance you used dsnow instead fod dS, and drain instead of dr

• We have changed the notation to be consistent (throughout the ms).

This definition is inaccurate. Lake desication starts at the point where E/I exceeds 1, in your case upwards of dss. d\* is the maximum isotopic enrichment that can be archived for open water bodies in an environment/setting. This is rarely achieved and if achieved it is for desiccating lakes with E/I>1. Desiccation starts earlier than d\*. If your argument is different please explain if not please re-define. One cannot enrich theoretically more than d\* because of the vertical isotope exchange from dA.

• By "desiccation", we more specifically meant "total desiccation", or the most enriched a water body may become under a given set of atmospheric conditions (air temp. and relative humidity). We have clarified this description to be "the maximum isotopic enrichment that can be achieved for a given set of environmental conditions" here and in the Figure 3 caption (L146).

use of dual isotopes to trace the composition of source water dates back to the work of Dincer, 1968 https://doi.org/10.1029/WR004i006p01289

• This is true, however the approach that we employ introduced by Yi et al. (2008) was the first to conserve the mass of both isotopes (hence "coupled isotope tracer method") when calculating E/I.

### correlation?

• We have changed to "correlation" here and elsewhere when appropriate (throughout manuscript). Can this be caused by lateral inflow of groundwater maintaining the lakes with small WAs? Please highlight if such deeper groundwater dependent lakes doesn't exist among the constellations.

• All land in this region is underlain by >100m of hydrologically impermeable permafrost that extends well beyond the bedrock, so no groundwater inputs to lakes exist beyond the subsurface runoff that occurs in the ~80cm soil layer between the permafrost and the soil surface (i.e. the active layer).

is this not that the LEL pass through the center and on average the number of samples plotting above and below the LEL must be equal. Unless otherwise there are outliers that impact the slope of the LEL?. The plot doesnt seem to also support the argument of above and below vs rain and snow. The LEL pointing to the rains is the best argument to confirm the influence or rains in the water balance. Alternatively, please indicate the number above or below

• We have not calculated the LEL using a linear regression through lake samples, but instead we have calculated the LEL independently of lake isotope composition, using deltaP and delta\* as the two anchor points. We then calculated lake-specific evaporation lines to calculate deltaI. This allows us to evaluate any tendency towards rain/snow sourced waters in the set of lakes, and also means the LEL will not divide the lake isotope compositions equally.

To display the number of lakes that have a more rainfall-like or snow-like deltaI by including deltaP on the y-axis of Figure 4a

Differences in dI across the sampling months is not discernible in figure 3 though figure 4a which i assume is drived from figure 4 (as in YI et al (2008) shows this. I leave this to the authors on how best they show this in figure 3.

• We chose to not plot dI Figure 3, since the points would cluster along the LMWL and overlap with precipitation and lake data, making it difficult to observe any patterns, hence why we only show dI in Figure 4a.

This is the key finding of the research. Linking lake isotopic composition not just to the physics of evaporation and water balance but to those watershed properties that govern or affect water balance. However, the key challenge is the applicability of such method in lakes with variable depth. Fortunately for the Thermokarst lakes the depth looks uniform from lake to lake. I wonder if one could reach at similar conclusions if the lakes did have variable depth in which case the lakes are fed by groundwaters or water residence time (V/I) wold impact the isotope enrichment. Thus the limitation of applying the method to lakes with variable bathmetry needs to be highlighted.

• We now address your comment about lake depth in the response to your review. Lake depth is mostly similar in this region to the lakes we sampled, but there are a few outlier lakes that are much deeper where the relationship we derived between WA/LA and Average E/I is likely less representative. We have added this discussion to the manuscript. (L328 – 334)

Reason? Could this be because of upstream lakes feeding the downstream lakes and thereby altering the isotope enrichment and E/I?

• Yes, we believe this is the reason and mention it in the discussion.

#### References

• Pienitz, R., Smol, J. P., and Lean, D. R.: Physical and chemical limnology of 59 lakes located between the southern Yukon and the Tuktoyaktuk Peninsula, Northwest Territories (Canada), 54, 330–346, https://doi.org/10.1139/f96-274, 1997.

#### **Reviewer 3**

• Replies to your comments in the PDF are in the attached PDF.

The authors investigated how lake and watershed properties and meteorological conditions influence the water balance of thermokarst lakes in the tundra uplands east of the Mackenzie Delta in the Northwest Territories of Canada. They sampled 25 lakes for isotope analysis five times in 2018, beginning before snowmelt on May 1 and ending on September 3 to calculate the ratio of evaporation-to-inflow (E/I) into the lakes and estimates the average isotope composition of lake source water ( $\delta I$ ). They indicated the range of average E/I ratios compared very well with watershed area to lake area (WA/LA) parameter (R2 = 0.74). Furthermore, they used this relationship to predict the average E/I ratio of 7340 lakes in the region, finding that lakes are not vulnerable to desiccation in the region. The authors have a good data and other measurements that could compliment isotopic computations to establish a surrogate.

While the general idea for finding a surrogate parameter which is easier and less expensive to measure or used to estimate E/I ratios laudable, the competent application of the time-tested evaporation model (water -isotope -mass balance method) used by the authors to justify this surrogate build up is doubtful. There are several assumptions that were made or glossed over in the paper that are difficult to justify, and these make it difficult to accept the results as they are presented. A major update of the assumptions and re-evaluation of the E/I computations is suggested.

• Thank you for your comments and suggestions on our manuscript. While we agree with many of your comments, we believe that you may have misinterpreted our approach to calculating E/I based on the presence of a singular LEL on Figure 3. The LEL on Figure 3 was placed there as a reference case only, and we have in fact calculated lake-specific evaporation lines for every lake water sample, as outlined in Appendix A. We agree with the suggestions you made under the assumption we had used a singular LEL approach, but we believe our actual methods address most of the concerns you had. We have attempted to clarify our isotope framework methodology in the revised manuscript by implementing your other suggestions (e.g., adding a table with all parameters used in the isotope framework).

Below are a few suggestions to improve the paper.

The authors indicated that the 25 lakes cover a range of watershed sizes, surface areas, among other characteristics (which include vegetation, land use type, rolling hills etc) and that the lakes were sampled as either a headwater lakes or a downstream lakes (line 70-75). The above information suggests the ff:

1) that the isotopic behaviour/ responses to evaporation from these lakes may vary widely and hence there is the need to individually and separately evaluate the LEL, h, delta P, delta A, and finally E/I and not lump them together as shown in Figure 3; I encourage the authors to study Bam and Ireson 2019 https://doi.org/10.1016/j.jhydrol.2018.12.032 since this work is similar.

- Although we display a LEL on Figure 3, we did not used a single evaporation line to calculate E/I for all lakes. We define the LEL (Local Evaporation Line), as being between δ<sub>P</sub> and δ<sup>\*</sup>, defined in the Figure 3 caption and in lines 118-119 from original submission "… *the LEL was anchored at* δ<sub>P</sub> and the theoretical isotope composition of a lake at the point of desiccation (δ<sup>\*</sup>)…" The LEL is used as a reference component in Figure 3, to show whether lakes tend towards rainfall- (above the LEL) or snow-sourced (below the LEL) waters.
- We have, in fact, treated every lake water sample individually when calculating δ<sub>I</sub> and E/I, by generating a lake-specific evaporation line for every lake water sample, i.e. calculating δ<sub>E</sub> and δ<sub>I</sub> separately for each lake water sample, as outlined in Appendix A, Lines 368-370 from original submission: "*The isotope composition of the lake-specific input water* (δ<sub>I</sub>) was calculated following Yi et al. (2008), where δ<sub>I</sub> is estimated as the intersection of the LMWL and the lake-specific LEL, defined as the line between the measured isotope composition of the lake (δ<sub>L</sub>) and δ<sub>E</sub>, which is the isotope composition of vapour evaporating from the lake.".
- We have used single values for h, δ<sub>P</sub>, and δ<sub>As</sub> because these lakes were all sampled in the same year, and are situated in a small enough region that it is reasonable to assume that the atmospheric conditions they experience are similar enough to warrant using a single value for h, δ<sub>P</sub> and δ<sub>As</sub>. We do see evidence there may be slight differences in δ<sub>P</sub> latitudinally as the distance to the ocean changes, however this should not affect our isotope framework greatly.

2) there is a need to account for the inflow and outflow fluxes to these lakes, I did not see this accounting process or equations, neither was there a stated assumption that these fluxes have been considered negligible within the case and the period and why;

It was not our goal to quantify individual lake water balance components (inflow, outflow, evaporation, precipitation) for each lake. We do not assume inflow and outflow to be negligible, but we assume that lakes remain at a relatively steady volume (inputs ≈ outputs) over the period that lake water isotope samples integrate a hydrological signal (water residence time), see Line 375 from original submission: "...assuming that lakes are well mixed and in hydrological and isotopic steady state." We believe it is reasonable to assume that lakes remain at roughly the same volume given that no lakes appear to experience desiccation and therefore are always fed enough water to remain above their sill level, maintaining a relatively stable water volume.

3) If even we assume that these lakes are well-mixed throughout the period (this assumption was not made though), precipitation addition and output fluxes are negligible and hence E/I ratios could be computed; we know a range of other factors that do significantly affect evaporation rates from lakes and hence lake isotope composition and ultimately the E/I. Among these are the exposed surface area, depth, the geometry/ basin bathymetry, vegetation cover etc., are all these the same of the 25 lakes? I believe the authors have sufficient data to account for these parameters, since the ultimate goal is to find a surrogate.

- As stated above, we do assume lakes to be well-mixed, and we do not assume outflow to be negligible, but we assume that the lakes are in a steady-state condition.
- Precipitation inputs are also integrated into the "inflow" portion of the Evaporation-to-inflow ratio, by nature of the isotope methodology.
- We have accounted for other parameters that have the ability to impact E/I and  $\delta_I$ . As described in lines 139-151 from original submission, we tested many lake and watershed attributes (all listed in Table 1) for significant correlation with E/I and  $\delta_I$ , however we only showed statistically significant relationships in Table 2. We clarify this point now in the Table 2 caption. "Only significant (p<0.001) relationships are shown in the table."

4) the choice or search for delta P (initial lake water isotope composition) using the intersection point of the LMWL and LEL is not clear. We see in Figure 3 that both May (and Sept ) samples plotted on or above the LMWL and if for anything the isotopic compositions of the May /June samples could be used as the starting point for those lakes to estimate the E/I between May and Sept or August. Extrapolating the LEL to the bottom to meet the LMWL to find initial source water has been proven to be unrealiable (see Bennettin et al., 2018, https://doi.org/10.5194/hess-22-2881-2018).

• We did not use  $\delta_P$  as the initial lake water isotope composition in our methods. We define  $\delta_P$  as the average isotope composition of precipitation, from which we anchor the LEL which we provide as a reference only. We calculated an "initial lake water isotope composition" (i.e. lake waters before evaporation took place) as  $\delta_I$ , as described in our response to comment #1. See Lines 368-370 from original submission: "*The isotope composition of the lake-specific input water* ( $\delta_I$ ) was calculated following Yi et al. (2008), where  $\delta_I$  is estimated as the intersection of the LMWL and the lake-specific LEL, defined as the line between the measured isotope composition of the lake ( $\delta_L$ ) and  $\delta_E$ , which is the isotope composition of vapour evaporating from the lake.". Since we are not computing a linear regression through lake water samples to define the lake evaporation line, our approach to calculating  $\delta_I$  is not vulnerable to the biases described by Bennettin et al. (2018).

I would recommend that since five (5) samples were collected from each of the 25 lakes from May to Sept, it will be useful an LEL is constructed for each of the lakes and used together with the rainfall isotopes and temperature measurements to estimate the dA for each of these lakes (see Bam and Ireson 2019; Bennett et al., 2008; Gibson et al., 2008a,b,2016; Gibson and Reid, 2014; Skrzypek et al., 2015). Use the May/June sample isotope composition as the initial lake water isotope to estimate the E/I for the period (May/June to Sept where feasible).

• The studies you have referenced seem to use variable  $\delta_A$  values because they use multiple years of water sampling, or because of large distances between lakes where one would expect differences in the atmospheric vapour source. Since all our samples are collected in the same year, and we assume that lakes experienced reasonably similar atmospheric conditions given their proximity to one another, we believe it is reasonable to use a single value for the isotope concentration of atmospheric vapour ( $\delta_{As}$ ).

There cannot be any E/I estimate outside the ice-free period, water fluxes lost during the dry winter periods originate from the sublimation of frozen ice over and snow, which covers on the surface of the lakes. The lakes are closed to evaporative losses prior to the open-water season.

• We do not claim that the water samples taken from lakes on May 1 represent the water balance conditions on that exact date, see L83-85 from original submission "*The May 1 samples also represent the hydrological status of lakes during the freeze-up period of 2017, since virtually no hydrological activity occurs during the winter months due to the complete freezing of the soil and lake surface.*". We use samples from May 1 because it allows us to evaluate how the freshet has influenced lake waters.

# PDF COMMENTS

chnages in long-term precipitation (since climate change would be attributed to a period not less than 30 years)

• This previous study actually found lakes responded to year-to-year changes in precipitation with no long-term trend. We have changed the text to reflect this. (L3)

in complete sentence; WA/LA has alreday been identified to be an inferential factor in lake water balance estimates

• This is already stated on L37-39 of the original submission: "Key drivers of lake water balances include the relative size of a lake within its watershed (Watershed Area/Lake Area, WA/LA), rainfall and snowfall patterns, permafrost dynamics, wildfire, vegetation cover, and ice-free season length (Turner et al., 2014; Arp et al., 2015; MacDonald et al., 2017; Wan et al., 2020)."

hmmm The paper cited has not been published neither is it accepted for publication - at this point it will be most useful to state how this was done.

• We have now added a brief sentence about how the correction was applied, and the paper is now published. (L92-94)

Same here, please state how these are done if that's something we need to know about

• As above.

only period that requires E/I ratio computations. Ref cited is incomplete since paper has not been accept for publication

• We disagree - it is valuable to use the samples taken on May 1 so we may see how the freshet period influenced lake waters in comparison to the ice-free period. The reference cited is now published.

It is interesting to note that there is an emphasizes on the range of watershed sizes but not on the range of the lake sizes. If I remember correctly, the geometry of lakes/ponds significantly influences the E (evaporation rate). Why make the this choice based on the range of watershed sizes?

• The words immediately following ("surface areas") refers to lake surface areas. We have emended the text to "lake surface areas". In Table 1, you will see that lake surface area ranges widely, from 0.37 to 90 ha. (L78)

What about compare how headwater lakes and the downstream lakes differ in E/I?

• We separate headwater and downstream lakes in Figure 5, and discuss the reasons for different E/I ratios between headwater and downstream lakes in lines 278-281 of the original submission.

before melt- at this stage only sublimation occurs; there is no evaporation of the lake water itself. The system is closed between Oct and May. Noting leave leaves and nothing enters

• We do not claim that water isotope samples do not reflect the lake water balance conditions for the specific date that the water samples were collected. The water isotope samples represent the lake water balance conditions integrated over the residence time of the water in the lake. We noted this in the following sentence of the original submission.

How did you account for these in and out fluxes from the lakes?

• We did not account for all components of the lake water balance individually - instead we use E/I ratios as a general indicator of lake water fluxes.

This seems to me to be the best time to executive E/I ratio computations and not before for the earlier reasons stated

• We use the 05-01 sample date in order to evaluate how the E/I ratio of the lake changed in response to snowmelt. As we in lines 83-84 and further down of the original submission, the 05-01 sample is representative of the lake at freeze-up in 2017 when lake inputs and outputs ceased.

where are the results or plot which indicate this ? What are the uncertainties associated with this?

• These results are shown later in the manuscript in the discussion section, and there we discuss some of the uncertainties associated with this approach.

I'm wondering what percentage this is, the Figure shows most lakes plotted closer to the rainfall region than the snow but clustered around the LEL indicating the strong influence of evaporation on the lakes

• We now include deltaP on the y-axis of Figure 4a to show the split between rainfall-dominated and snowmelt-dominated lakes for each sampling date.

What do you mean, here? Not clear- lake water isotope compositions will surely differ due various climatic and physical factors but what do you mean by along the LEL, d18OI and E/I? Each lake must give you a distinct LEL, d18OI and perhaps E/I. How did you compute these?

- We are stating that we observe changes in d18OI and E/I between the sampling dates. We removed "along the LEL" since it is somewhat redundant in the sentence. (L206) We have improved our explanation about how the LEL, d18oI and E/I are computed in the methods section (L154 -166), and added in more detail in Appendix A.
- We now include deltaP on the y-axis of Figure 4a to show the split between rainfall-dominated and snowmelt-dominated lakes for each sampling date.

The colors used in this plot are confusing; for example, snow is depicted as black circle and the same applied to the average rainfall and the limiting isotopic composition dssL and d\*. Is this plot for an individual lake overtime or different lakes with single samples overtime? If these are different lakes with signle samples over time, it will very difficult to use this a s justification for any E/I ratio computations. The authors have already indicated in line 70 -75 under study area how these lakes differ in various terms; headwater vs downstream, surface area exposure vrs size , veegattion cover : shrub vs tall tree etc all of these produces different effects on evaporation and isotopic composition hence these lakes cannot just be lumped together to make E/I estimates. Now as you can see from Figure 3 even the Sept samples are more depleted than the May samples and some of the May samples and Sept samples have plotted on the LMWL. If this is the case, then it is untenable to use delta P as your initial input isotopic compositions. There is difficulty estimating the initial isotope using extend LEL and LMWL intersections see Benettin et al 2018 https://hess.copernicus.org/articles/22/2881/2018/

• We have clarified that these are isotope values for all lakes for all sampling dates in the figure caption. We also added a legend for clarity. -We have changed all symbols on this figure are now all unique to the type of isotope data they represent. -As stated in our other comments, we did not use deltaP in our E/I calculations. Individual deltaI estimates for each lake water sample were used in calculating E/I. The LEL is plotted on the figure only as a reference for interpreting the shifts in lake water isotope composition.

What do you mean by recharge phase? Does groundwater vertical infiltration (recharge) cease because the surface of the lakes are frozen during the winter? Not sure this a correct assumption

• The Recharge Phase is described in the previous paragraph, it is the period when inflow is greater than evaporation and causes E/I ratios to reduce. However, groundwater infiltration does cease at some point in winter because the entire soil column freezes in winter. Only larger streams may have any unfrozen sediments below their bed, however there is usually little lateral movement.

I would recommend you make a list of all the abbreviated parameters used in the model to the difficulty encountered in reading the text.

• we have added a table to the Appendix that lists and describes all the parameters used in the isotope framework and how they are generated.

References

Skrzypek, G., Mydłowski, A., Dogramaci, S., Hedley, P., Gibson, J. J., and Grierson, P. F.: Estimation of evaporative loss based on the stable isotope composition of water using Hydrocalculator, 523, 781–789, https://doi.org/10.1016/j.jhydrol.2015.02.010, 2015.