# **DSM2** Bay-Delta Tutorial 3: Source Tracking (Fingerprinting)

**Purpose:** The purpose of this tutorial is to use the source tracking capabilities of the model to create a fingerprinting study. We will set up both volumetric and concentration-based fingerprinting and visualize the results.

# 1. Reopen the historical tutorial

- a. In windows, navigate to \{DSM2\_home}\tutorial\historical.
- b. In the GUI, open historical\_tutorial.

### 2. Create a model for source tracking:

In the background, source tracking imposes a computational cost on QUAL that is the same as one additional constituent per source. For this reason, it is useful to comment out source tracking.

- a. In *historical\_qual\_ec.inp*, locate the GROUPS include section.
- b. Uncomment the group definitions for source tracking. You may wish to review this file to see how the groups are identified.
- c. Similarly uncomment the two finger printing files the ones that have "source\_track" in their names.

## 3. Define volumetric inputs

a. Create a file called tutorial\_volumetric\_fingerprint.inp. Go through each of the time series input files for QUAL and create an equivalent input that has a constant value of 100 with the constituent called volume. This step is conceptually simple, but will produce a large file – feel free to break it into several files if you prefer.

#### 4. Define the fingerprinting output

a. Specify Clifton Court concentration output for each of the source groups that you defined for both EC and for *volume*. The name should be clifton\_court, the concentration should be ec or volume and the interval should be 1day. Avoid

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redundancy -- you do not need to put the constituent name or the source into the output name: ie, use "clifton\_court" for the name, not "clifton\_ag" or "clifton\_ec"

5. Run HYDRO and QUAL for One Year

a. Change the model name in qual\_ec.inp and run HYDRO and QUAL for one year in 2002. Start QUAL a day later to avoid mass conservation errors in the first hour. Make sure the init\_conc variable is set to zero so that there will be no initial condition contribution for any variables (note: for a volumetric fingerprint, it may be useful to make this concentration 100 if you want to include initial conditions in the fingerprint analysis).

### 6. Process the output

a. Use VISTA or HEC-DSSVUE to open up the output file. Copy May-September concentrations for each location. Paste the output into a new sheet in the Excel provided called excel\_fingerprint.xls, which you can use as a reference. Use the "stacked area plot" in Excel (one of the standard Excel plot types) to plot up the fingerprint results.

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