

ECO Lab

Exercise 1: Modelling E.coli bacteria



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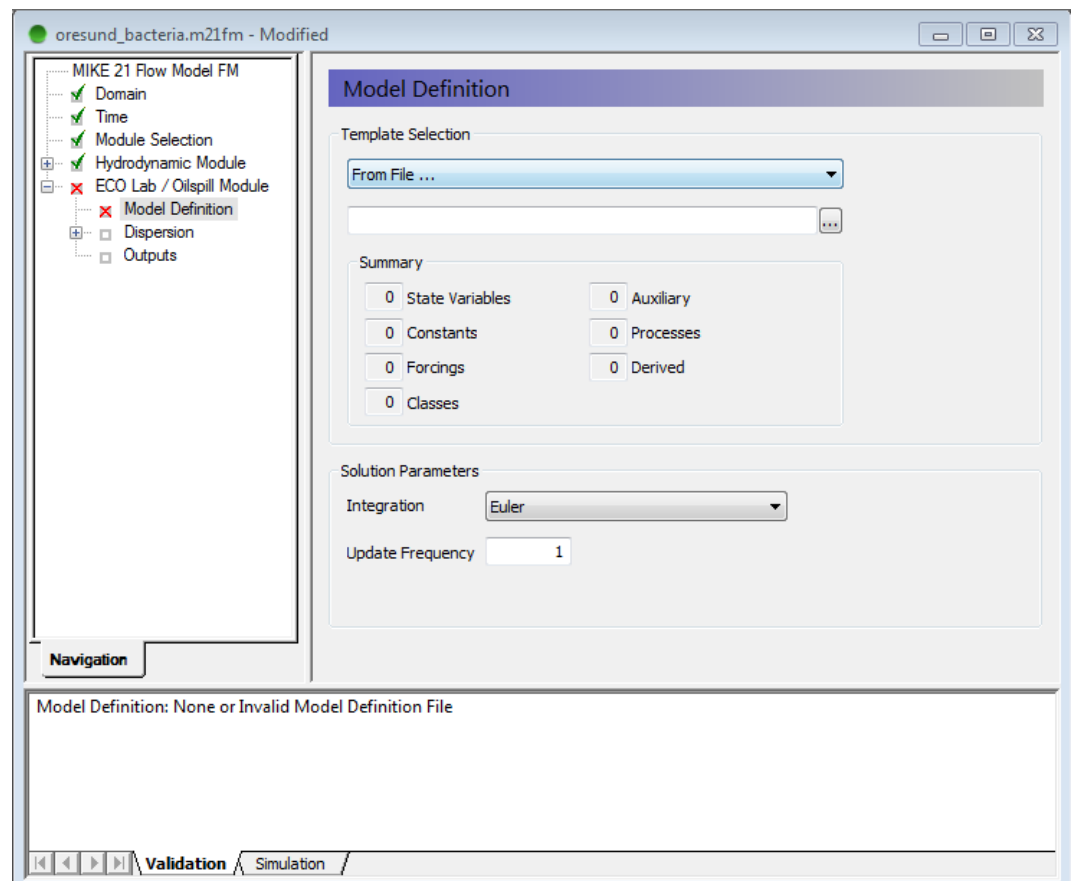
1 Exercise 1: Modelling E.coli bacteria

Getting started

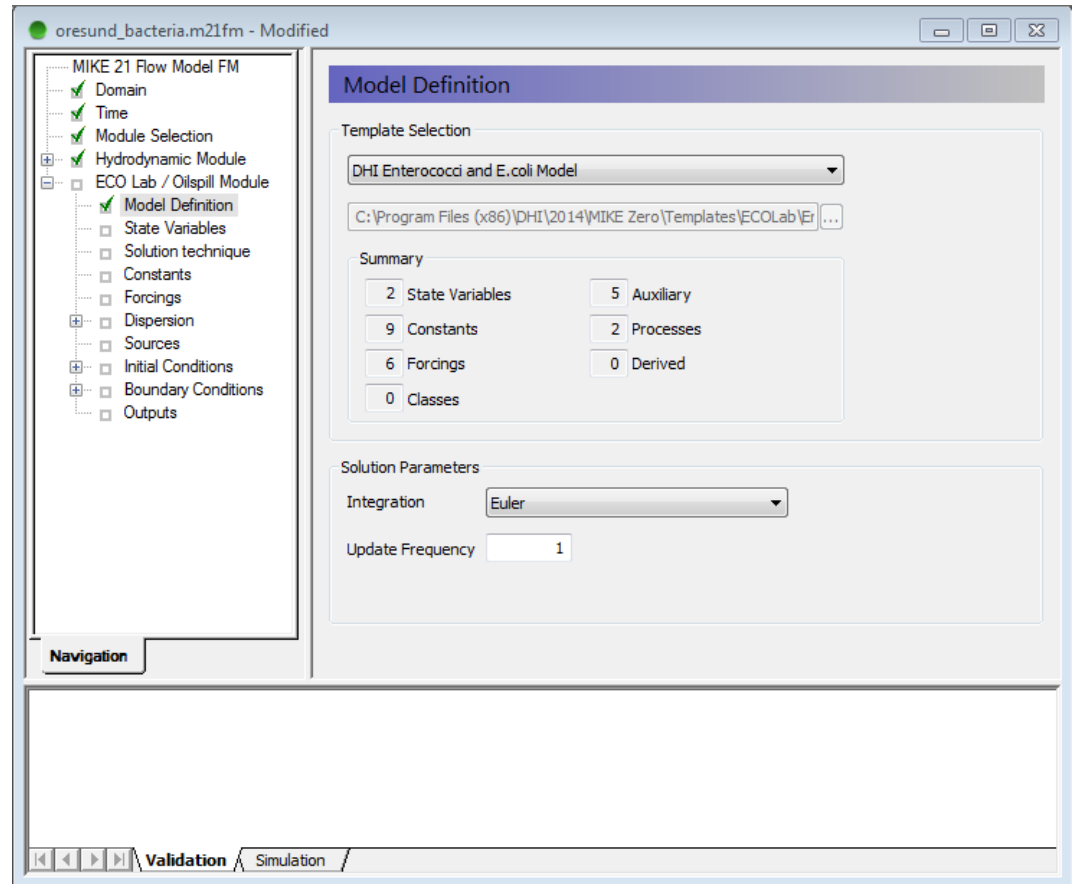
1. Locate the example folder:
Explore the folder C:\Program Files (x86)\DHI.... and look for ...MIKE Zero\Examples\MIKE_21\FlowModel_FM\HD\Oresund\Calibration_2
2. Open the file: oresund.m21fm
3. Save the file into a working folder and change the name of the setup file.

Template selection

4. In module selection, include ECO Lab.
5. In the model definition you open the 'From file ...' menu.



6. Choose the DHI Enterococci and E.coli model.



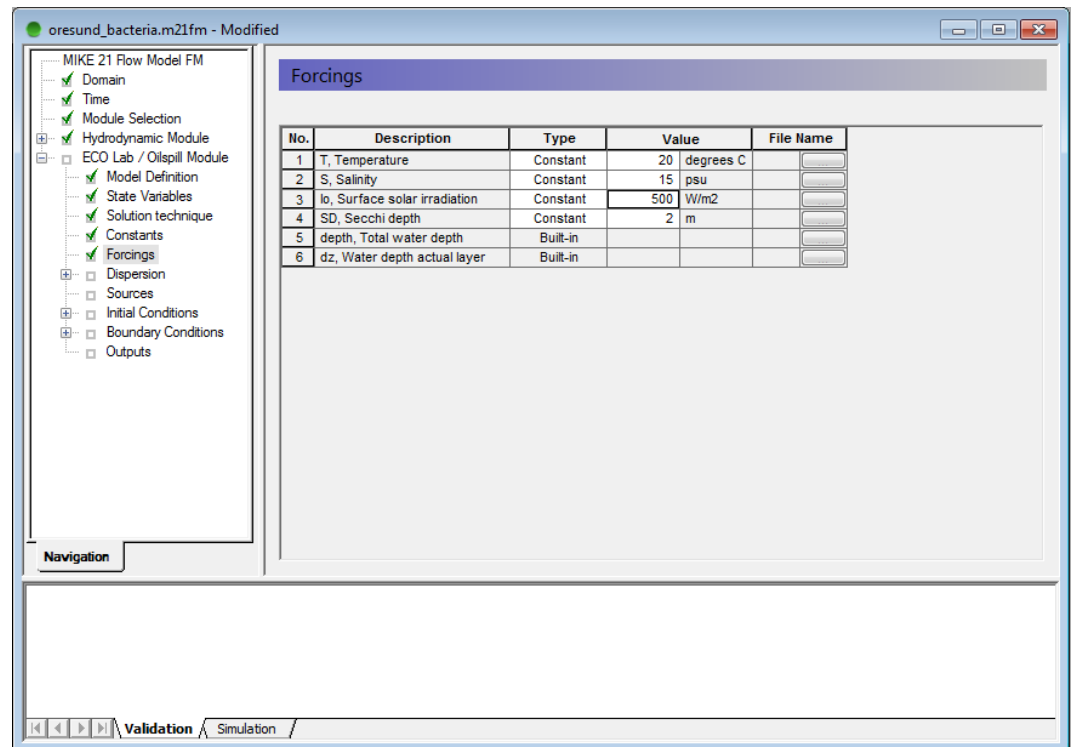
7. Set the update frequency to 5.

State Variables, Solution Technique and Constants

- For this exercise you do not change anything in the menus: State Variables, Solution Technique and Constants.

Forcings

- In the 'Forcing menu you include the constants as stated in the power point.



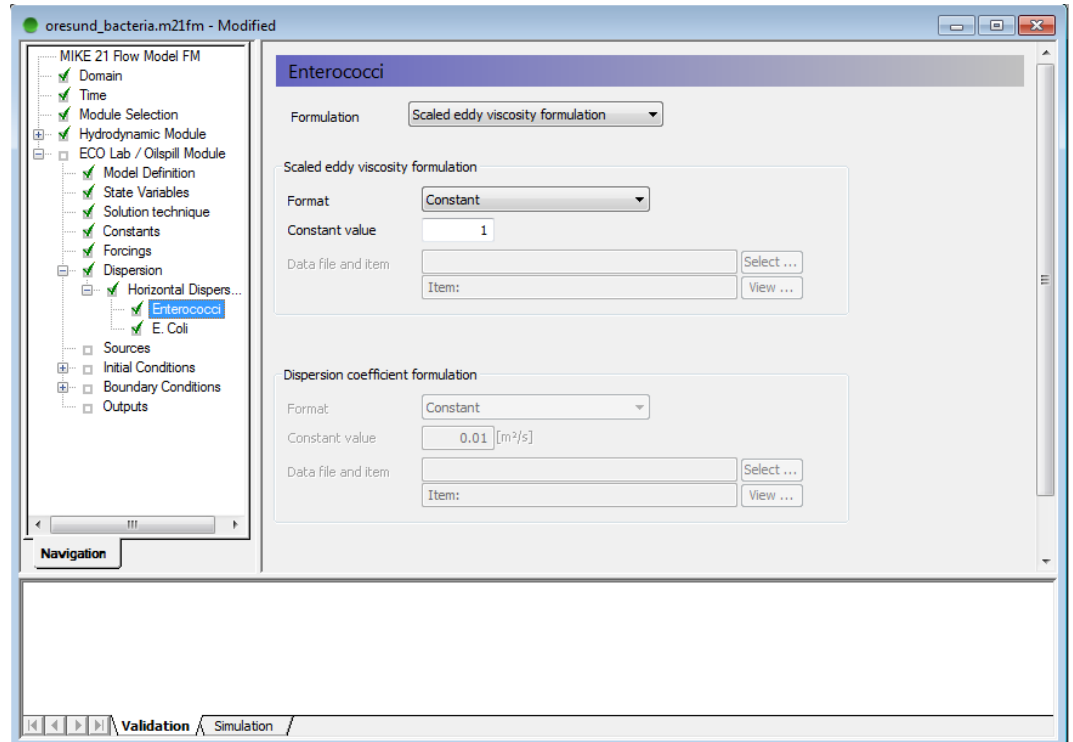
The screenshot shows the MIKE 21 Flow Model FM interface. On the left is a navigation tree with the following items checked: Domain, Time, Module Selection, Hydrodynamic Module, ECO Lab / Oilspill Module, Model Definition, State Variables, Solution technique, Constants, and Forcings. On the right, the 'Forcings' menu is open, displaying a table with the following data:

No.	Description	Type	Value	File Name
1	T, Temperature	Constant	20 degrees C	
2	S, Salinity	Constant	15 psu	
3	Io, Surface solar irradiation	Constant	500 W/m2	
4	SD, Secchi depth	Constant	2 m	
5	depth, Total water depth	Built-in		
6	dz, Water depth actual layer	Built-in		

At the bottom of the window, the status bar shows 'Validation' and 'Simulation' options.

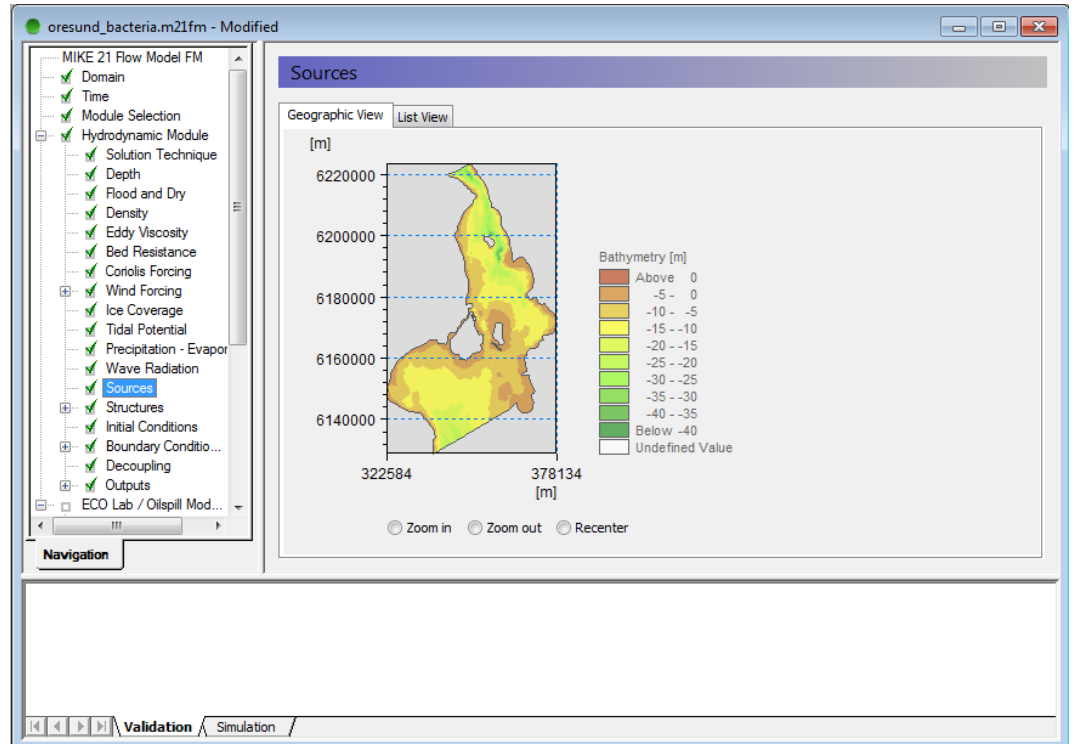
Dispersion

- For the Dispersion you include 'Scaled Eddy Viscosity Formulation' and keep the constant value at 1.

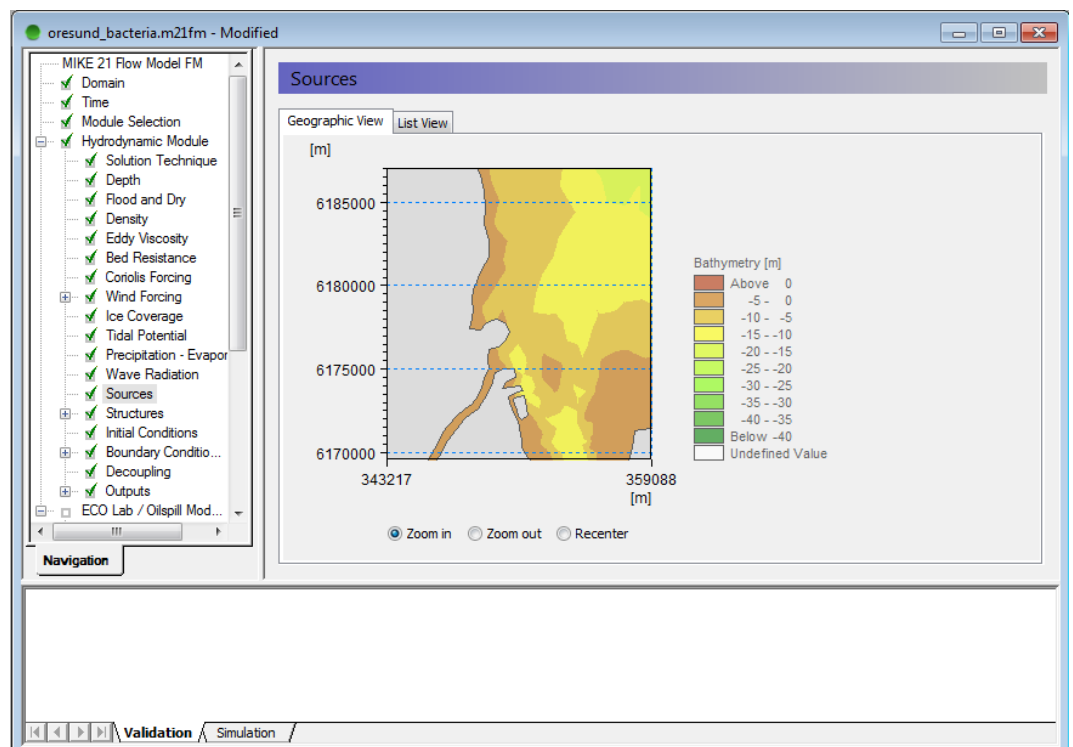


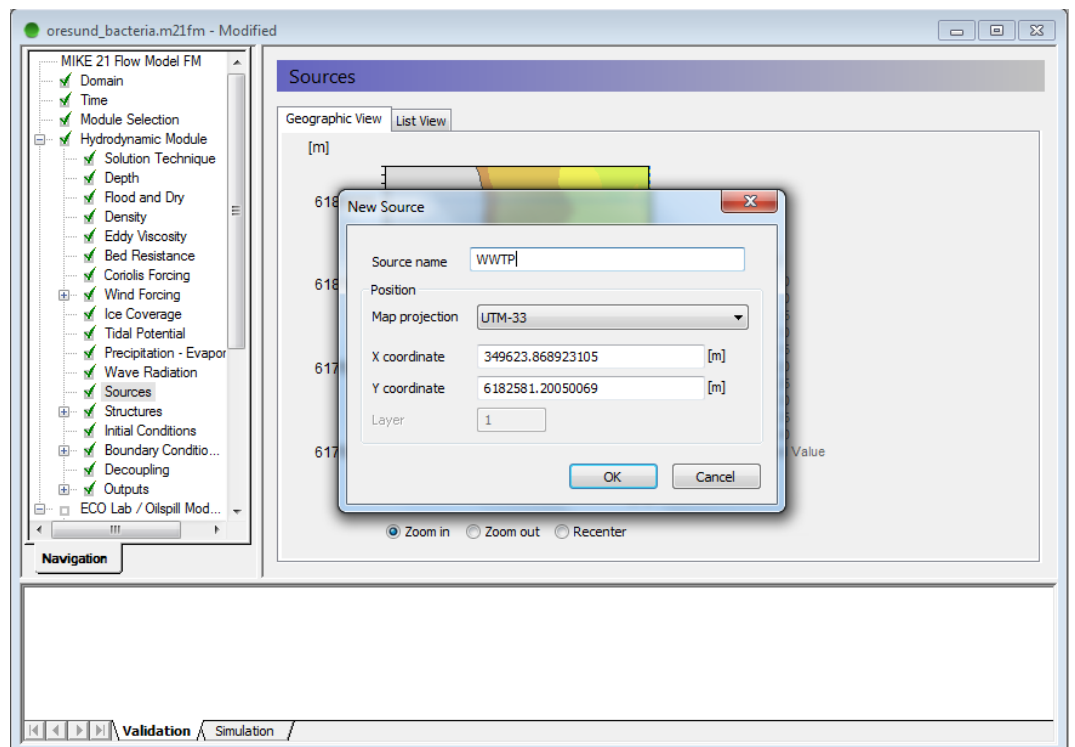
Sources specification

- For the sources you first need to include one source in the 'Hydrodynamic section' – open the hydrodynamic module and enter the 'Source' folder.

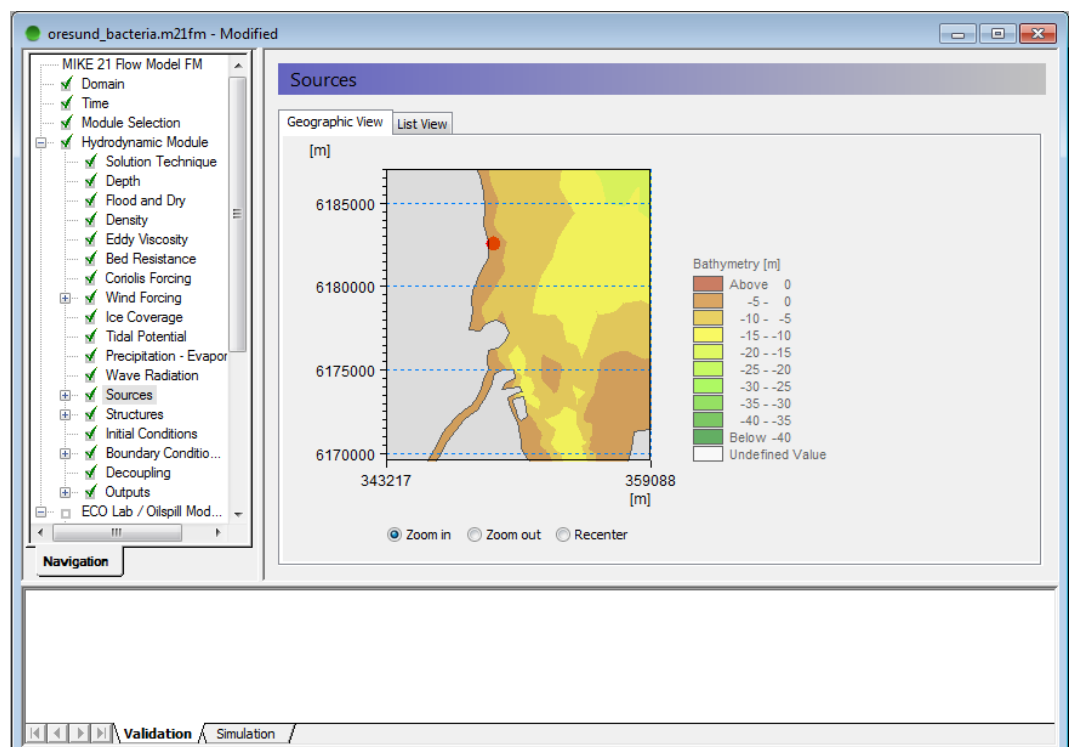


- Zoom in to a coastal area and double click where you would like to include the source.

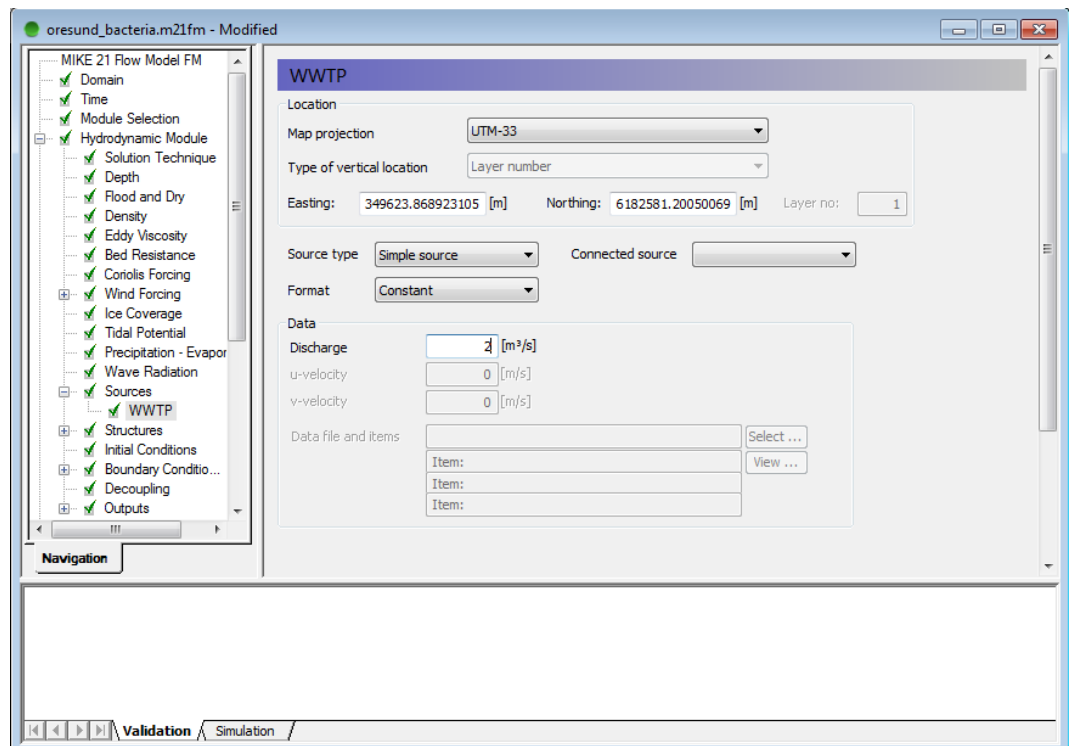




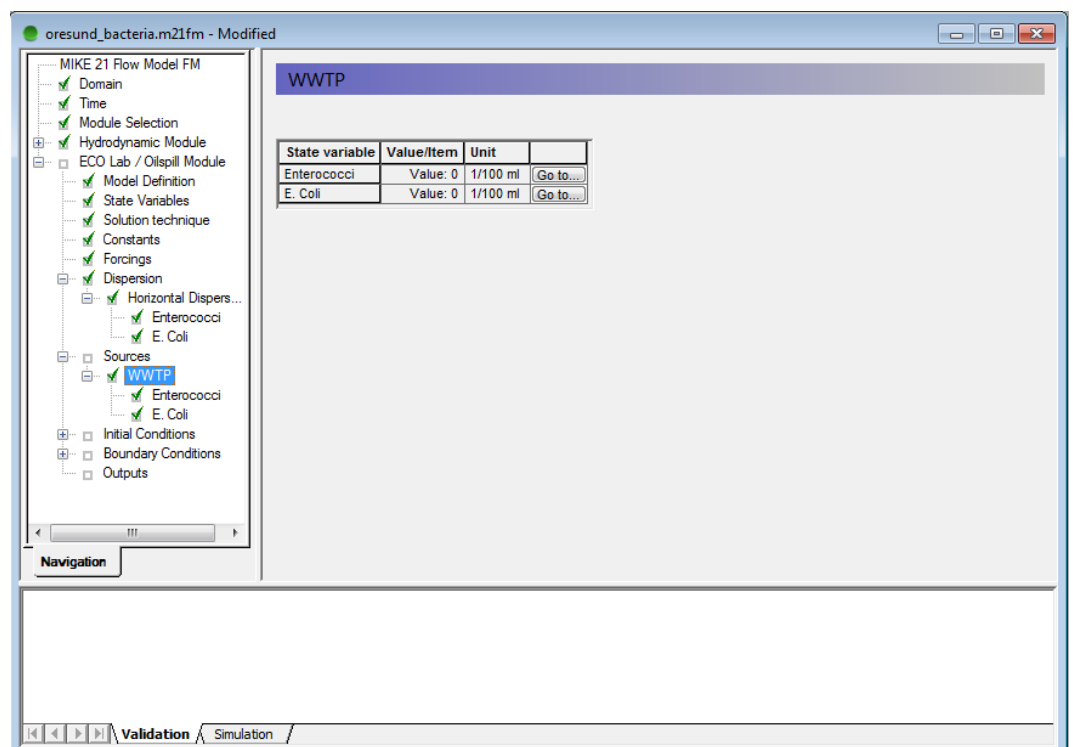
13. You now have a source, and the location is shown on the map.



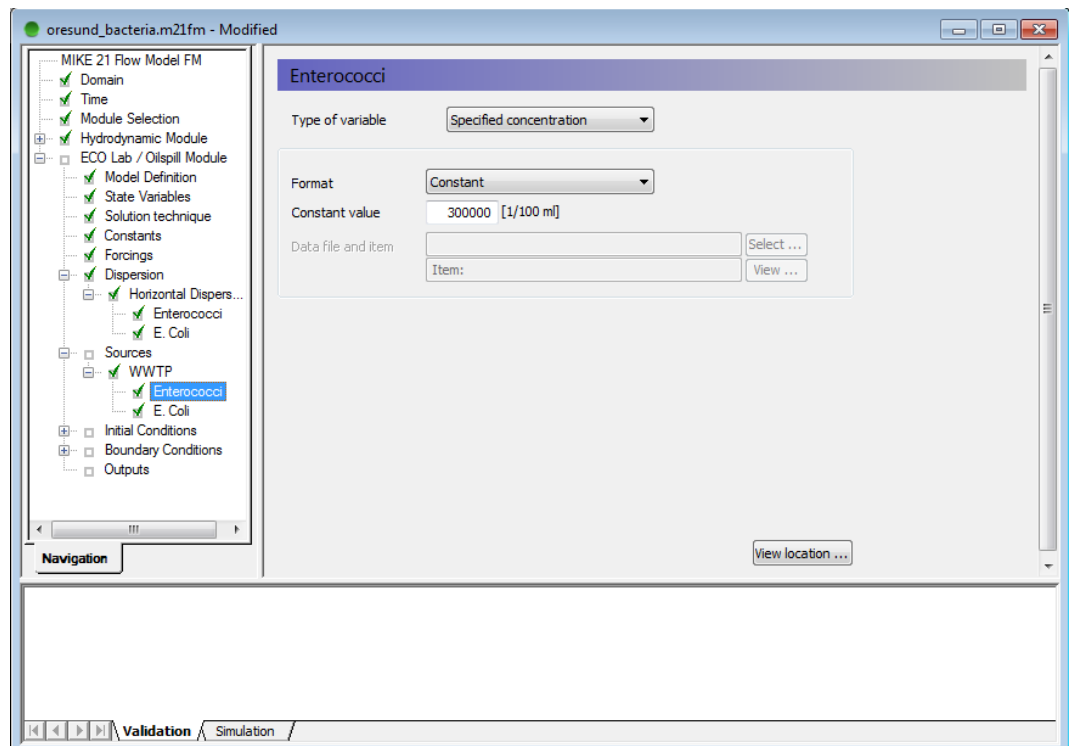
14. Enter the list view, press the 'go to' button and include a discharge as listed in the power point.



15. Close the 'Hydrodynamic section' and enter the 'Source' menu in the ECO Lab section.



16. Press the 'Go to' button and enter concentrations as listed in the power point.
300000 enterococci/100 ml and 1000000 E.coli/100 ml.

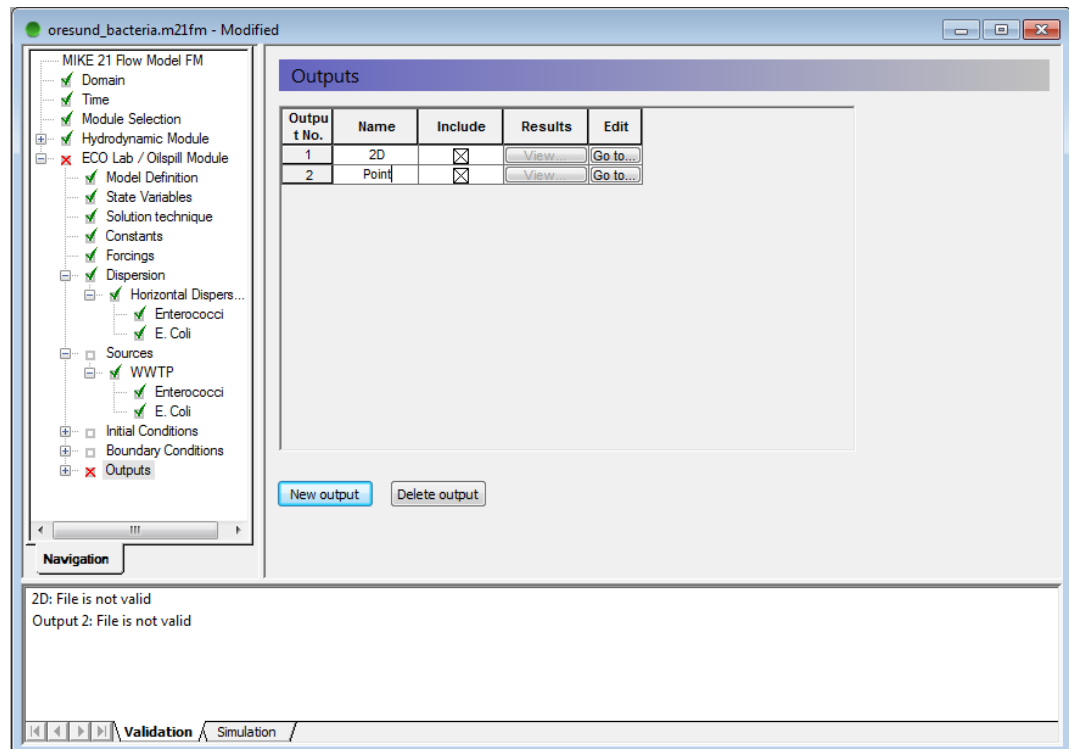


Initial Conditions and Boundary Conditions

17. Default values for the two bacteria are zero, why we do not need to change anything in the menus for 'Initial Conditions' and 'Boundary Conditions'.

Output specification

18. Enter the output menu and include two new outputs. Name them 2D and Point.



19. Create a 2D output file and a similar time series output as shown in the figures below.

Point: File is not valid

	Easting	Northing	Layer no.	Name
1	322028.77431	6127731.4889		
2	322028.77431	6224518.3037		
3	378689.6147	6224518.3037		
4	378689.6147	6127731.4889		

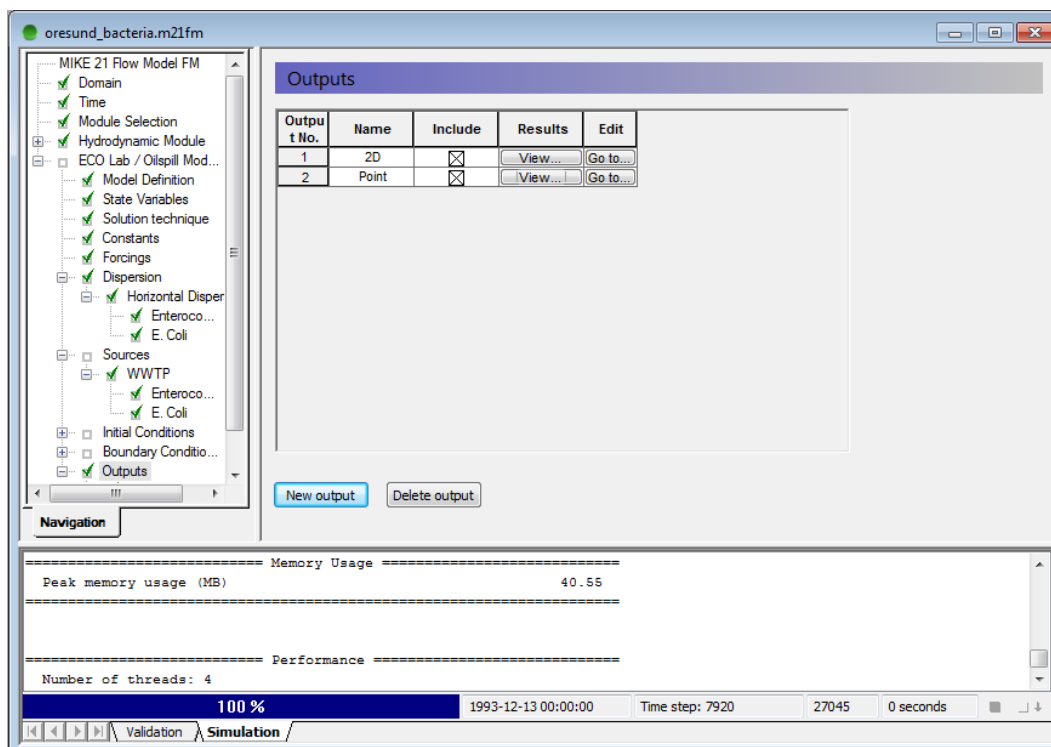
	Easting	Northing	z	Name
1	350000	6182000		Point 1

Model execution

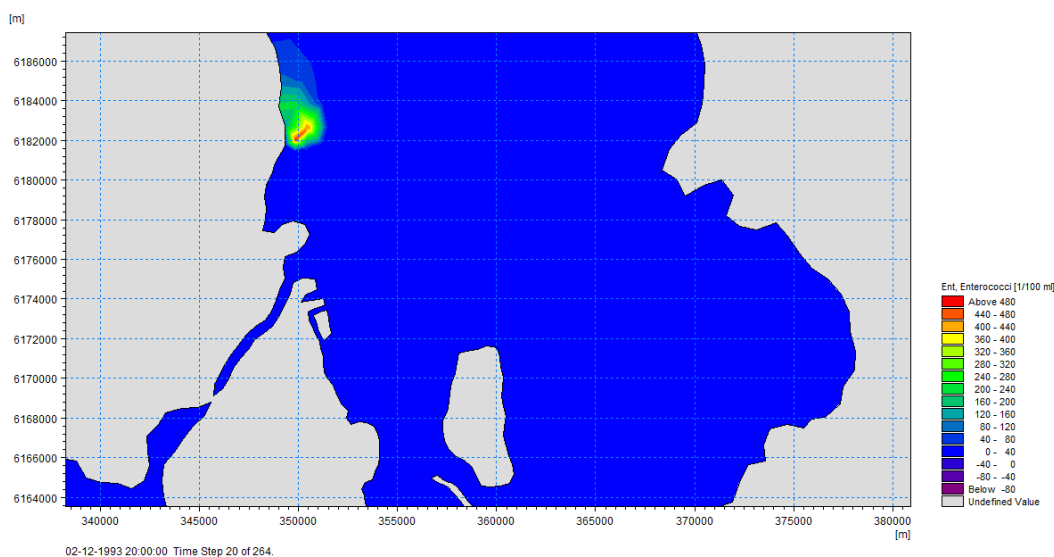
20. Run the model. If the model does not start running, please remove the output from the 'Hydrodynamic selection' (the folder for the HD results might be 'read only').

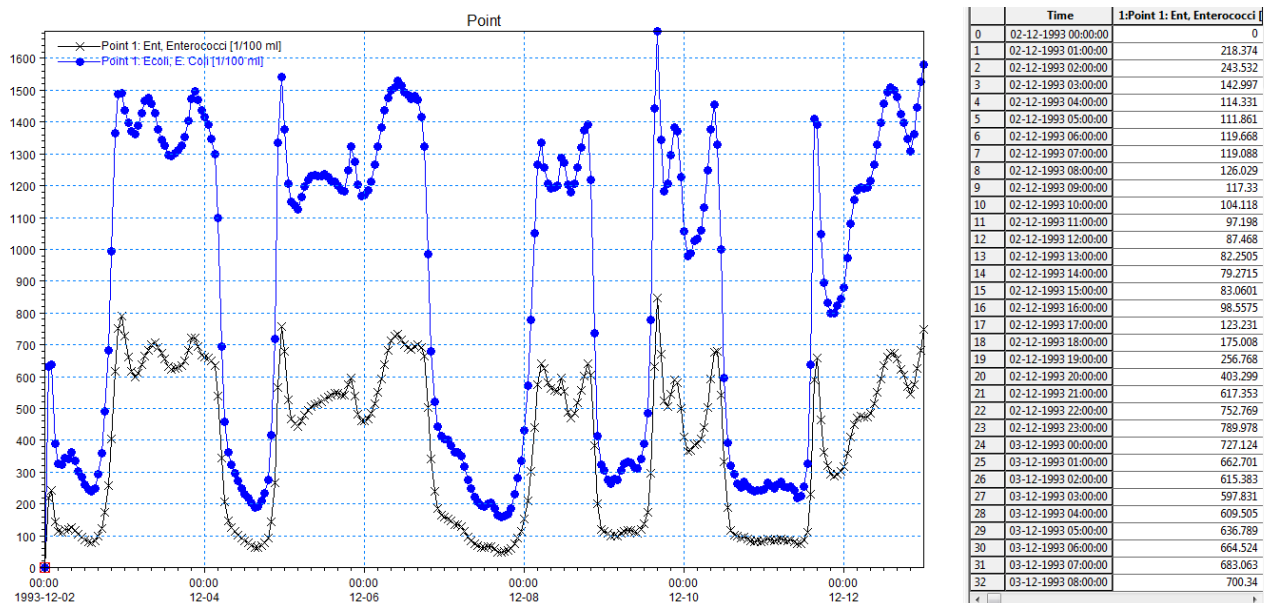
Model results

21. When the model run is completed the model results can be viewed from the output menu, by pressing the 'view' button.



22. The 2D results provide an overview of the transport and dilution of the bacteria source, whereas the point series show the results over time. The variance in the concentrations is governed by the current direction.





23. Applying the statistics from the 'tool' menu you can calculate the average concentration from the two bacteria: Enterococci and E.coli. For the example provided the average of the two are 379 enterococci/100 ml and 876 E.coli/100 ml.
24. Now try to change the Secchi depth in the 'Forcing' menu from 2 m to 5 m. Save the setup in a new name and repeat the model run.
25. The statistics from the model results now show 277 enterococci/100 ml and 575 E.coli/100 ml. Similar tests can be executed by changing temperature, salinity etc.
26. As stated in the power point, you can also try to do the same exercise by applying a 3D model setup. In this case include the source in the surface layer, but keep the remaining constants as in this 2D example.