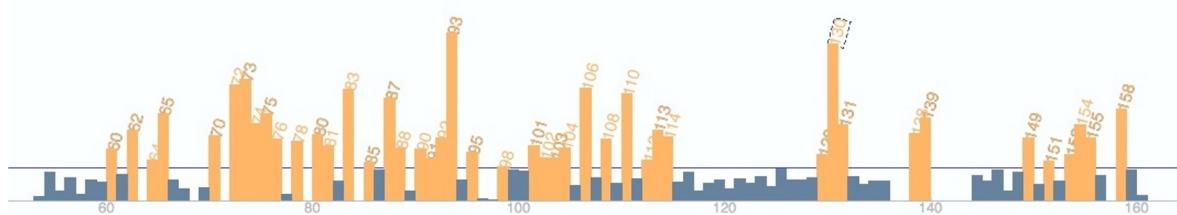
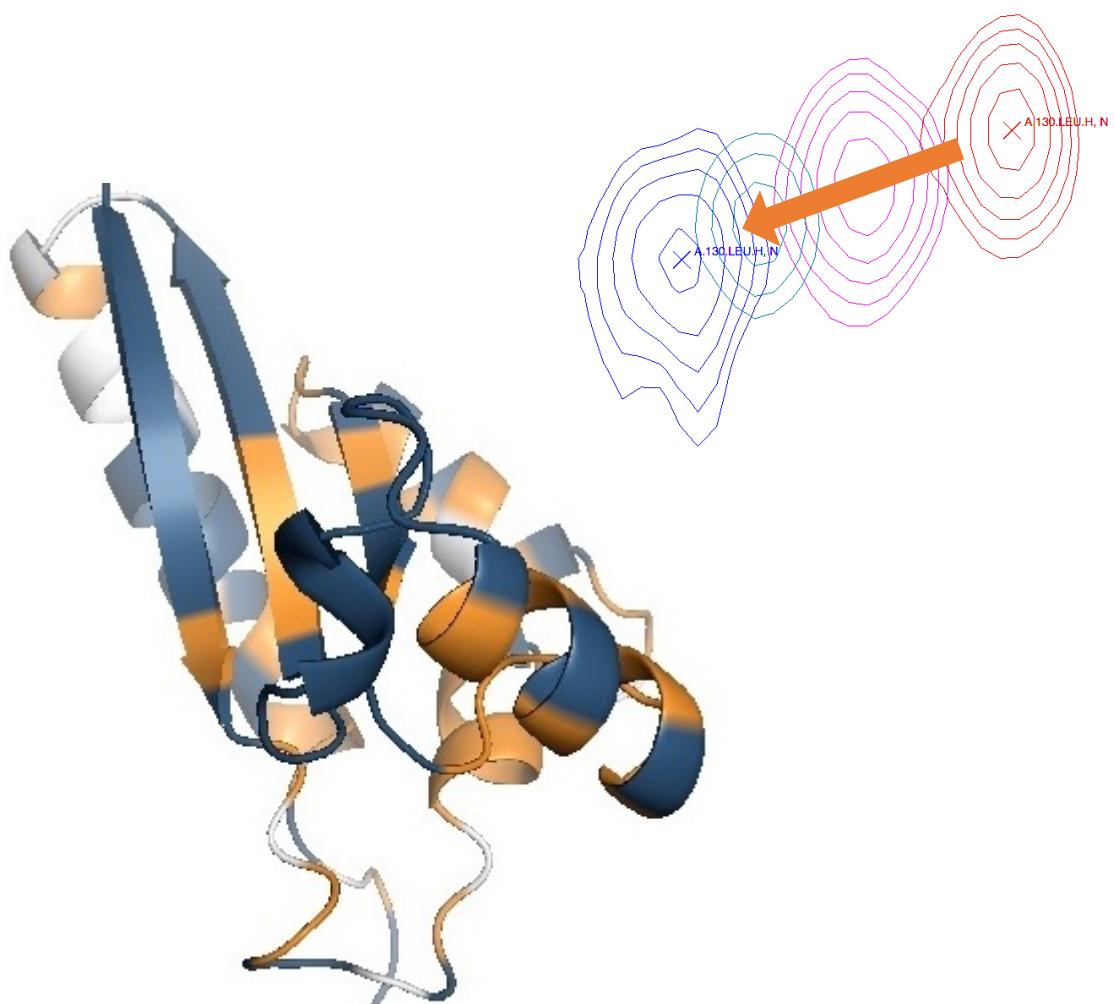


Chemical Shift Perturbation Tutorial



Introduction

This tutorial is designed to show you how to use CcpNmr Analysis Version 3.1 to analyse the chemical shift perturbations caused by a protein interacting with a binding partner (in this case an RNA molecule) and map the interaction onto the structure of the protein.

The tutorial uses ^1H - ^{15}N HSQC spectra of ^{15}N labelled TSTAR protein in complex with RNA. We are grateful to Dr Cyril Dominguez for making this data available to us. The original paper describing this work is [Feracci, M et al. \(2016\) Nature Communications 7, 20355](#).

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1. Open Project
2. Copy Peak List
3. Move Peaks
4. Map Peaks
5. PyMOL
6. Binding Curves
7. Issues

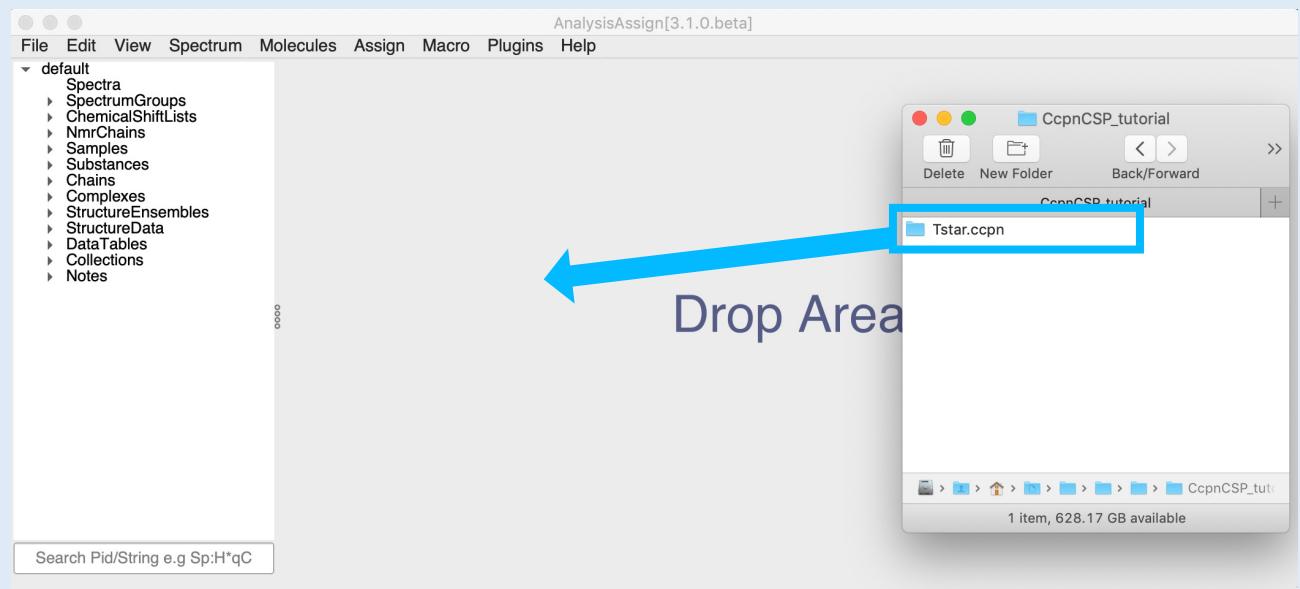
Start CcpNmr Analysis V3

Apple users by double clicking the icon  *CcpNmrAnalysis*

Linux users by using the terminal command: *bin/assign*

Windows users by double-clicking on the *assign.bat* file

Open Project



1A Drag & drop “Tstar ccpn” into the sidebar or Drop Area.

CcpNmr projects have an extension of type “filename.ccpn”. Find the project file *Tstar.ccpn* in the top directory or in AnalysisV3/data.

- Select the file “*Tstar.ccpn*”, drag and drop into the program. The Tstar project will be loaded in a new window.

You will see five spectra, displayed as:

Tstar-free (red)

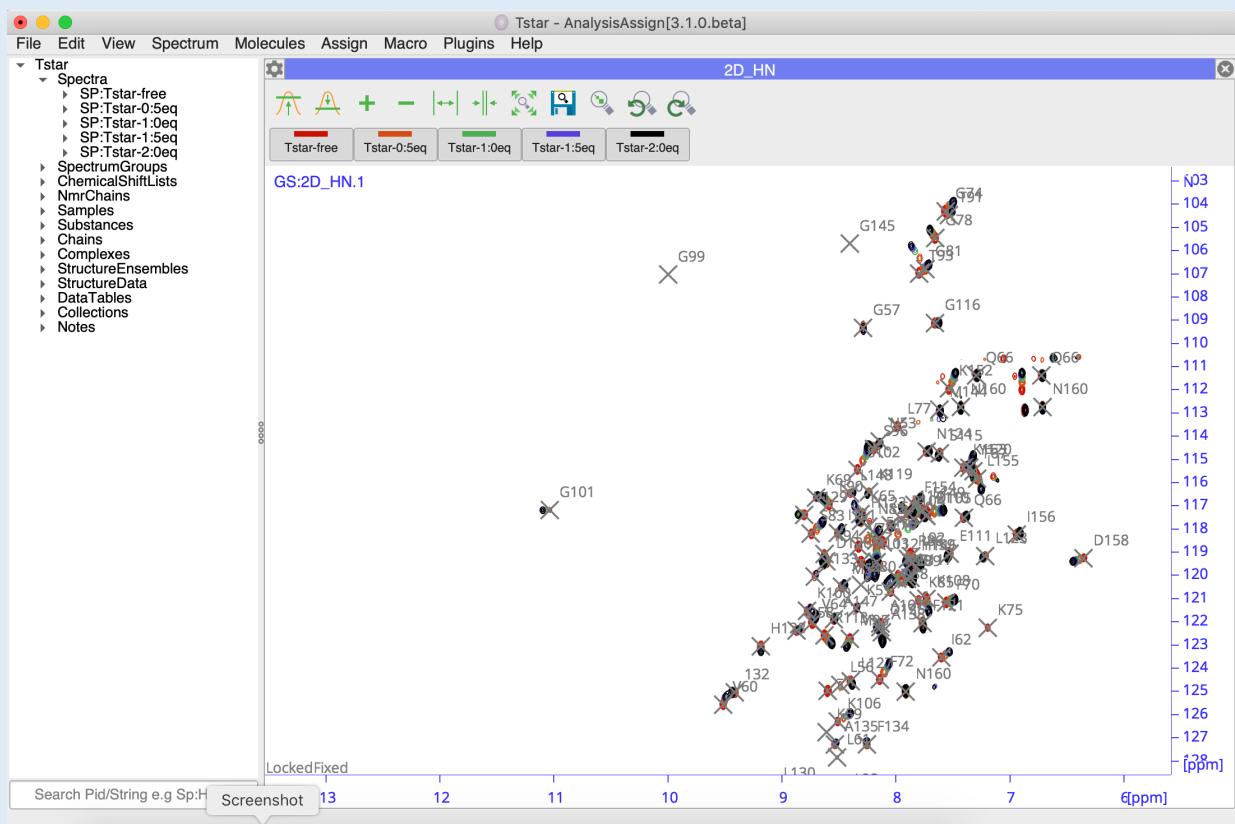
Tstar-0:50eq (orange)

Tstar-1:00eq (green)

Tstar-1:50eq (blue)

Tstar-2:00eq (black)

Open Project



Getting started, basic operations

Sidebar

All spectra and peak lists are located in sidebar. Double click on an item will open the properties popup.

Display

A display can contain multiple overlaid spectra. To show/hide a single spectrum, click on its toolbar button. See next page.

Mouse

- Pan → Left Click and drag in display
- Zoom in/out → Scroll wheel in display
- Context menu → Right click
- Select a peak → Left click on a peak symbol “X”
- Move a peak → select first then hold Middle or Right and drag

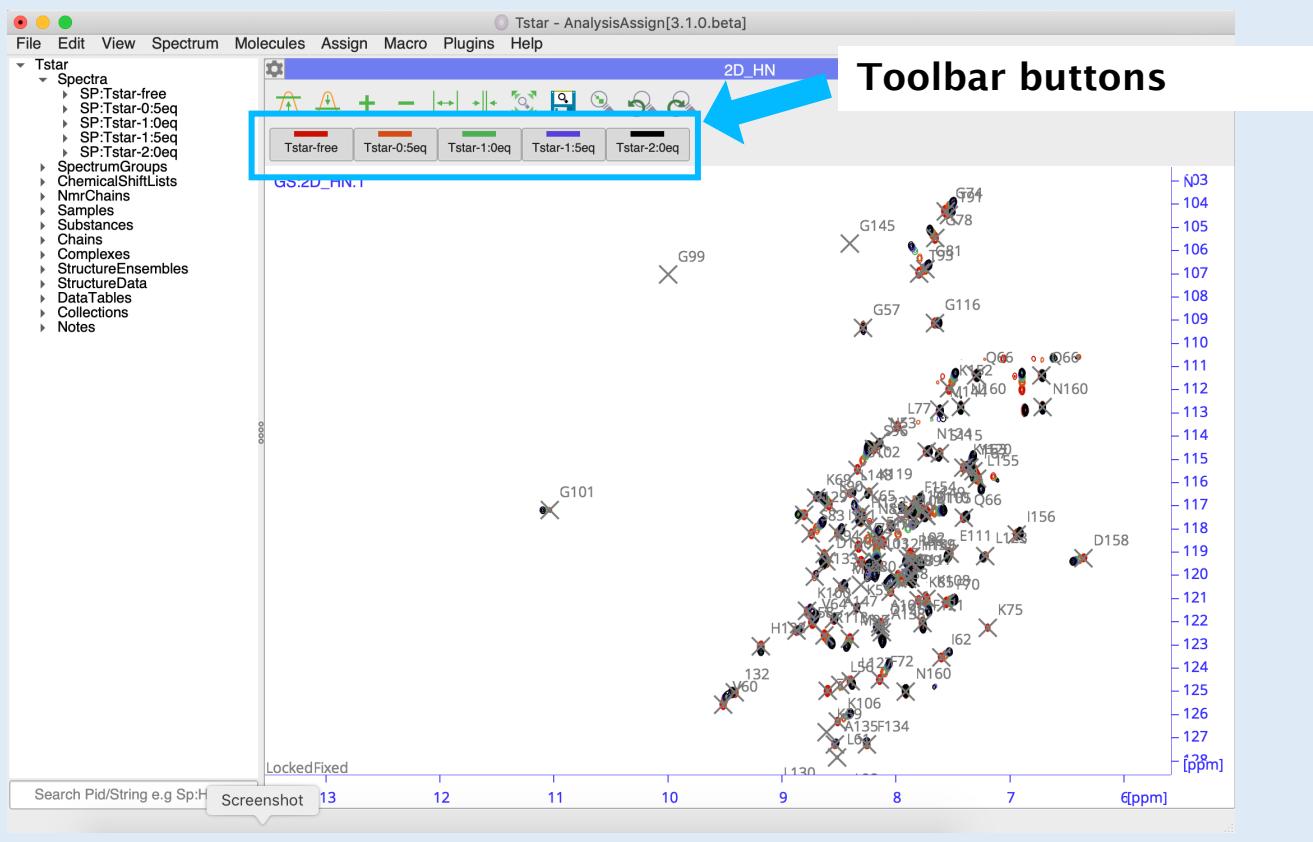
Shortcuts

The program uses several shortcuts, example “CL” for copying a peak list. You will need to press the first letter on your keyboard e.g. “C”, followed by the second letter, e.g. “L” .

For more commands and operations:

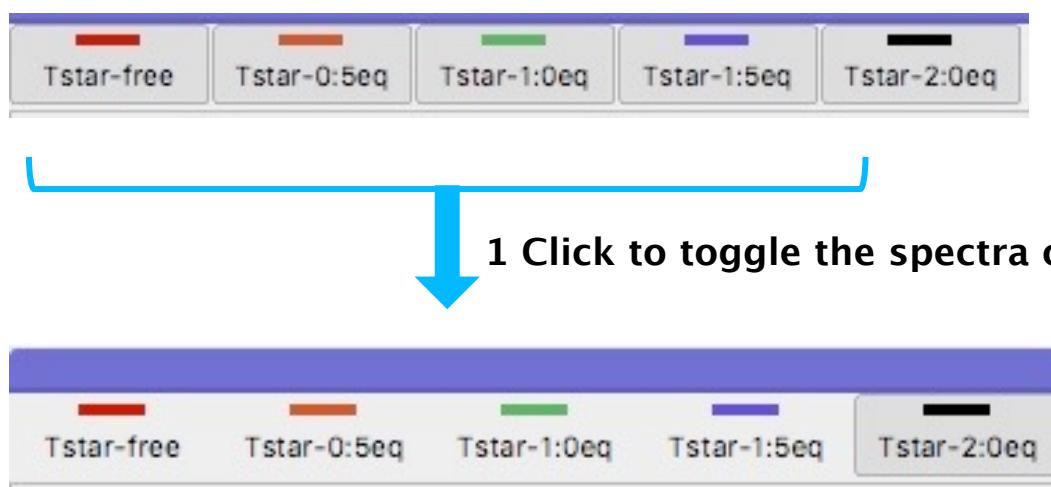
main menu → *help* → *Tutorial (Beginners)* or *Show Shortcuts*

Open Project

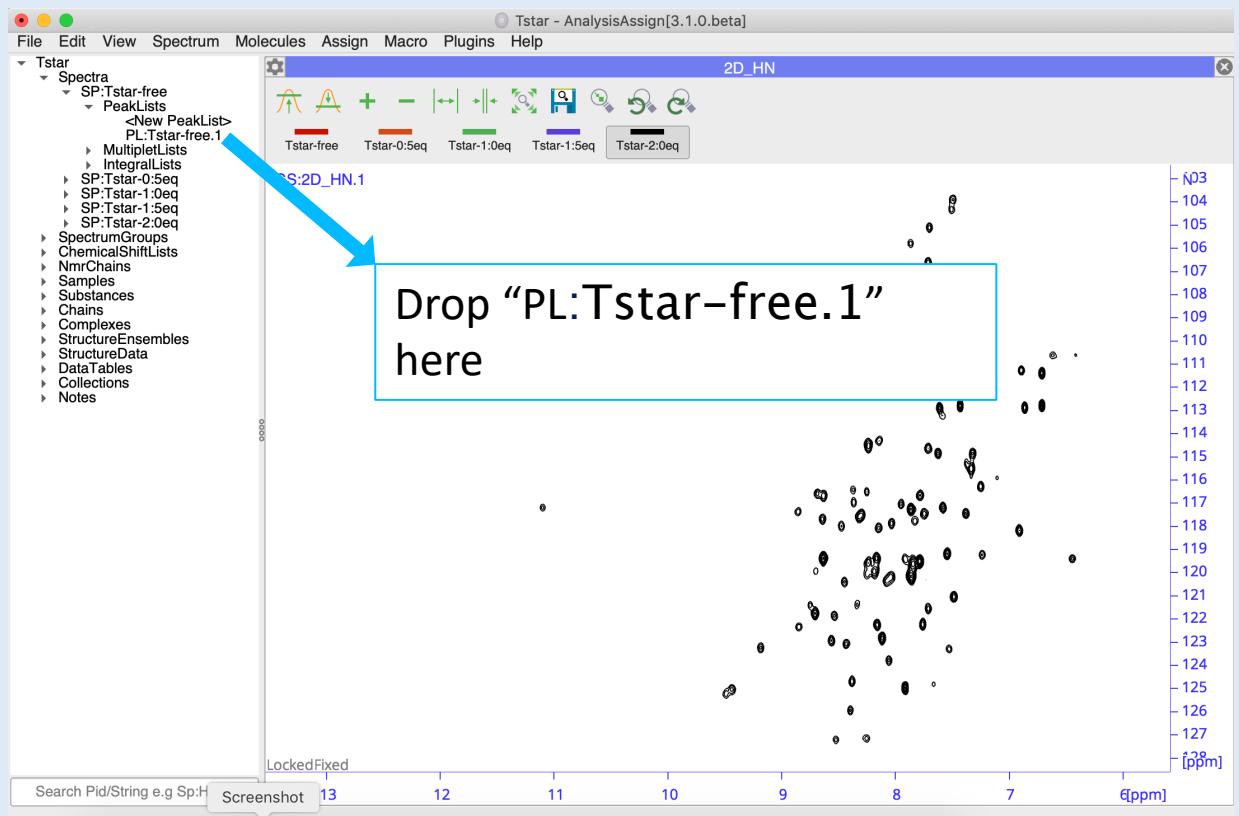


1B Toggle off all spectra apart from Tstar-2:00eq on the toolbar.

- Click on the toolbar button of the first four spectra (*a, b, c, d*).
The buttons will change on (spectrum visible on the display, “toggle On”) to off (spectrum hidden on the display, “toggle Off”)



Copy Peak List



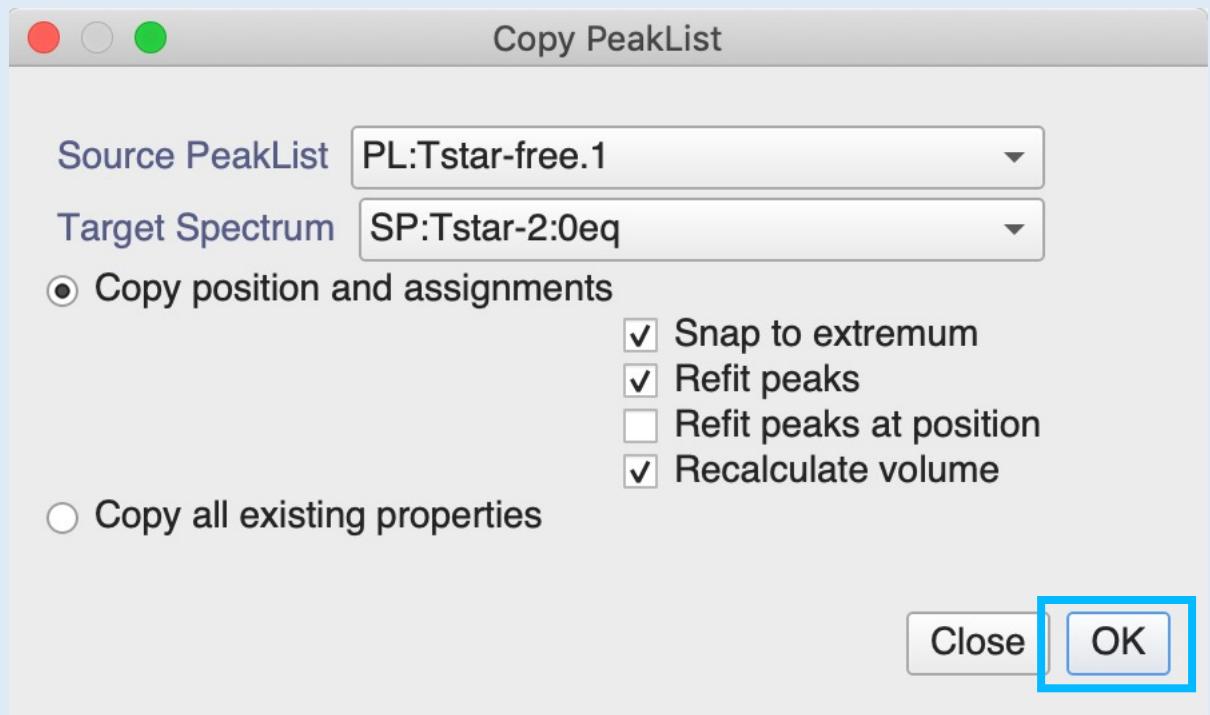
Only the Free-Form Tstar has a peak list. You will need to copy this peak list to a spectrum recorded with Tstar bound to the RNA.

2A Drag & drop the peak list tree item “PL:Tstar-free” into the spectrum display.

- On side bar, expand the tree:
 - ▼ Tstar
 - ▼ Spectra
 - ▼ SP:Tstar-free
 - PL:Tstar-free.1**

- Click to select PL:Tstar-free.3, drag and drop the item into the spectrum display.

Copy Peak List



2_B Select target Spectrum: Tstar-2:0eq.

A new popup will open, the first entry is the peak list source you want to copy; the second entry is the target spectrum, where the peak list will be copied.

Select:

- *Source PeakList:* PL: Tstar-free.1
- *Target Spectrum:* SP:Tstar-2:0eq

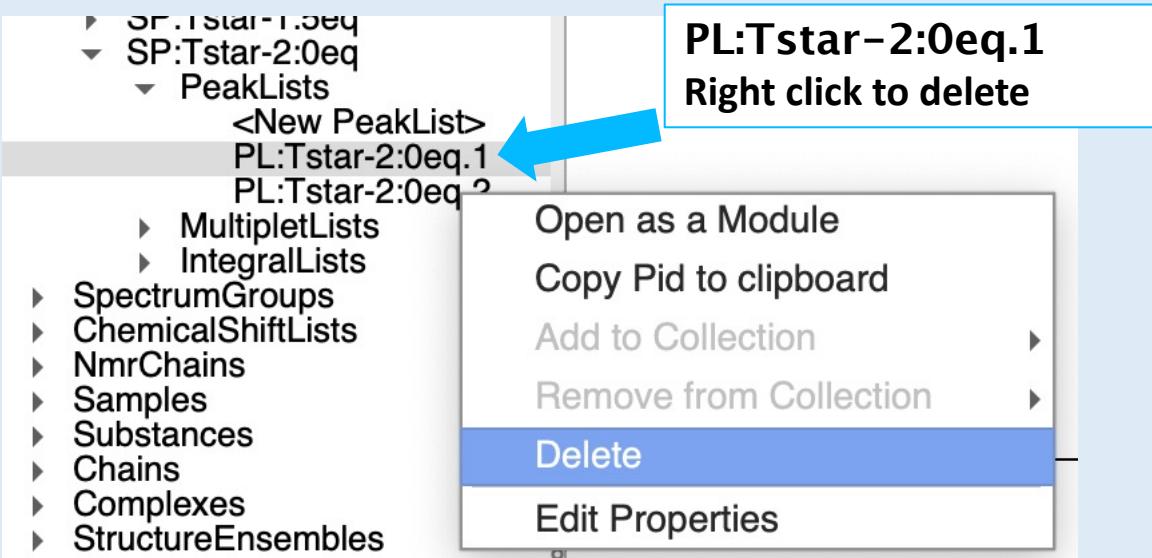
Leave the rest of the settings as they are.

- Press Ok

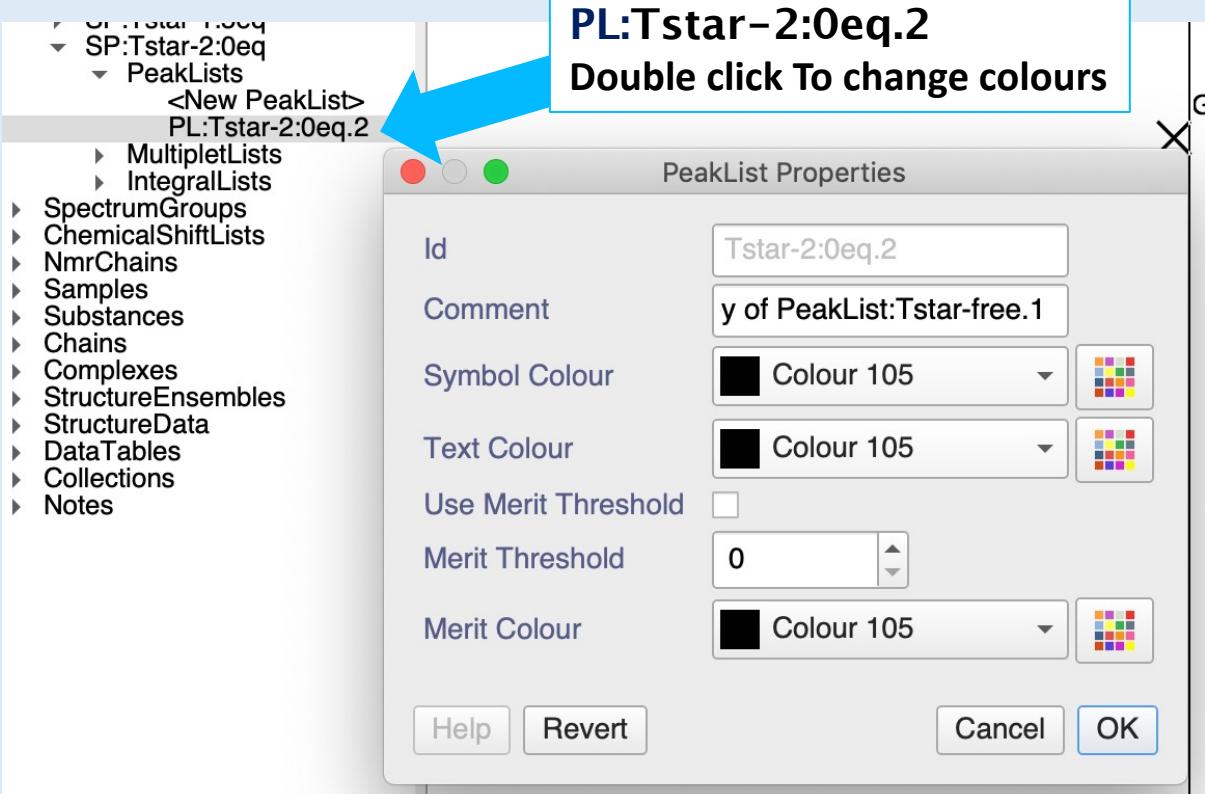
2

Copy Peak List (Optional)

2C



2D



2C

Delete “PL:Tstar-2:0eq.1” from side bar.

The default peak list for the spectrum Tstar-2:0eq is empty. For simplicity delete it from side bar. Right-click on the item **Tstar-2:0eq.1** then click delete.

2D

Double Click on sidebar item “PL:Tstar-2:0eq.2 ” to change colours.

- Expand the spectrum item tree *SP:Tstar-2:0eq* on the side bar, as previously for *SP:free-tStar*
- Double click on **PL:Tstar-2:0eq.2**

A new popup will open.

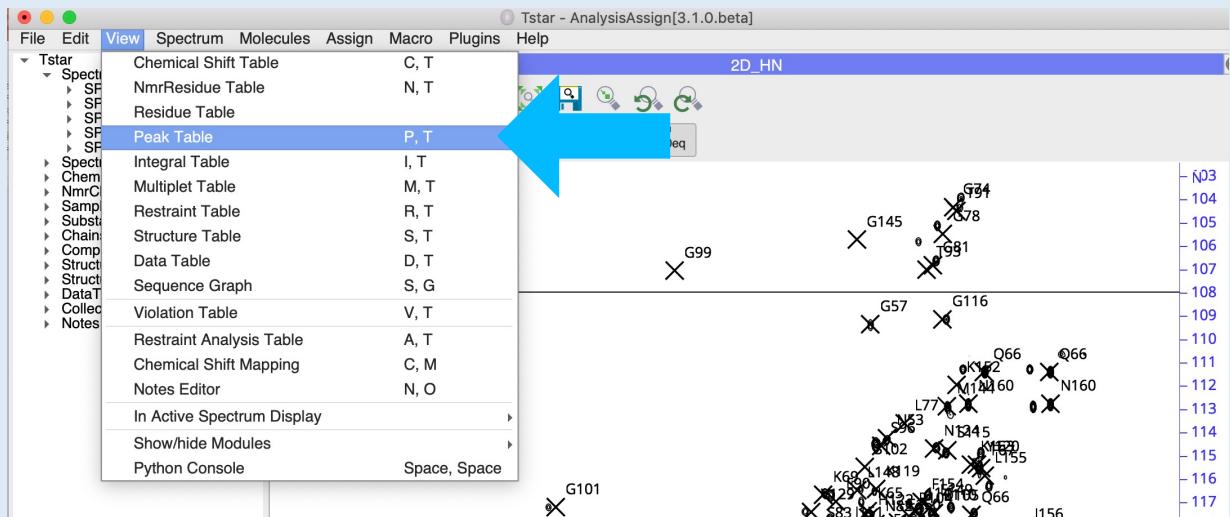
- Select the desired colours for text labels and symbols. Close the popup.

Tip: use the same colour as the spectrum contours.

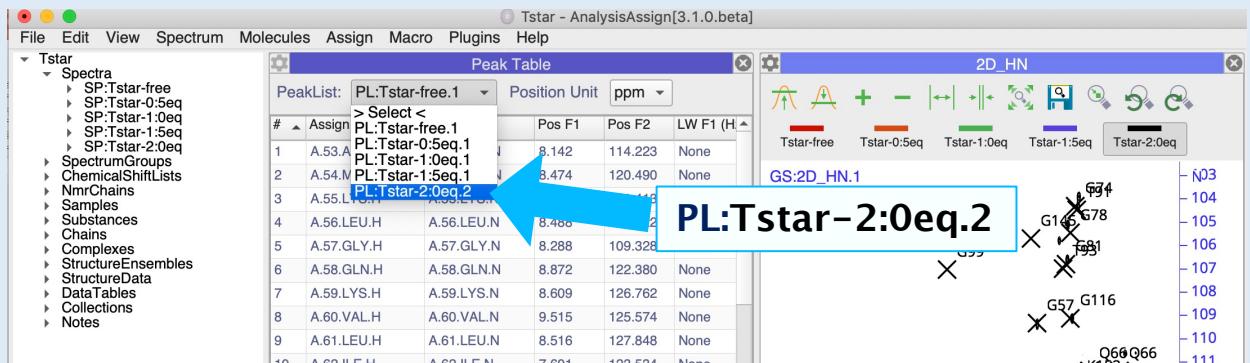
2E

Repeat steps 2A to 2D for each spectra.

Move Peaks



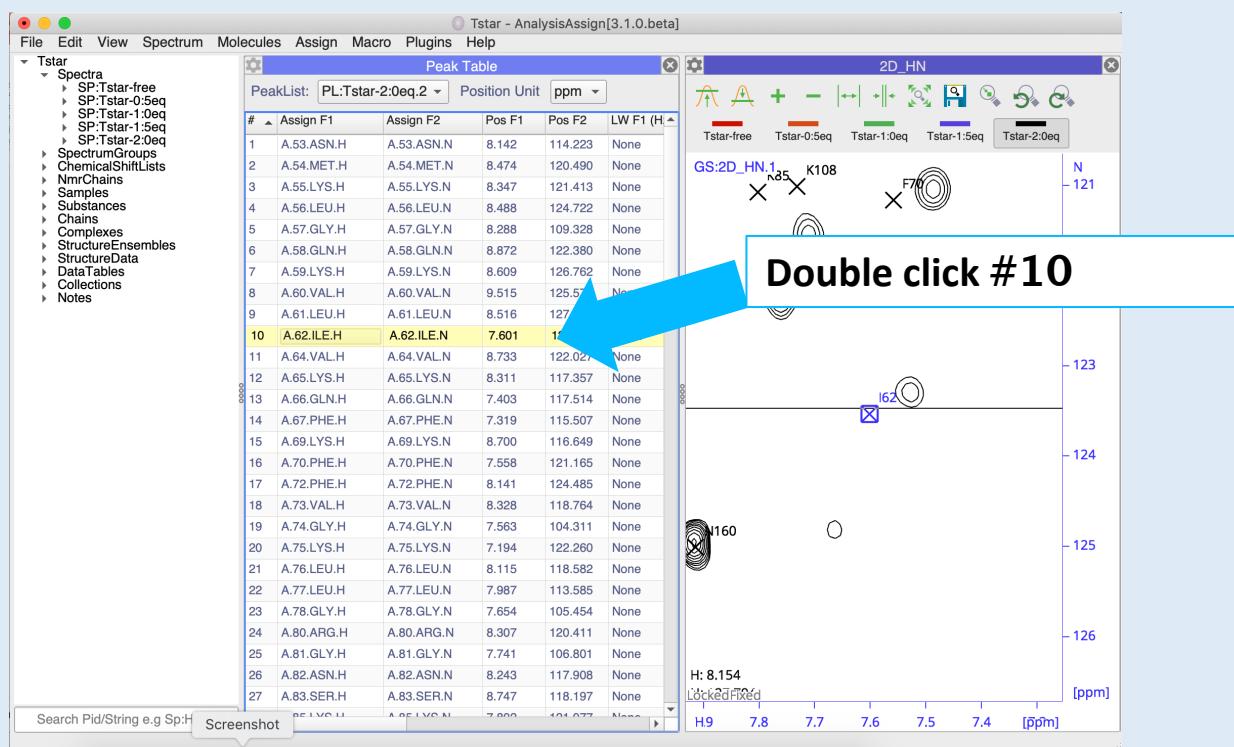
3A Open peak table, shortcut “PT”. Open a peak table with the shortcut “PT” or from the main menu –> *View* –> *Peak Table*.



3B Select PL:Tstar-2:0eq.2.

On the top left corner of the peak table there is a pulldown menu, select the newly created peak list: PL:Tstar-2:0eq.2.

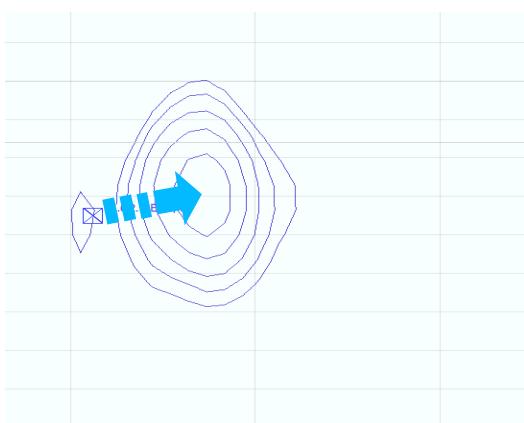
Move Peaks



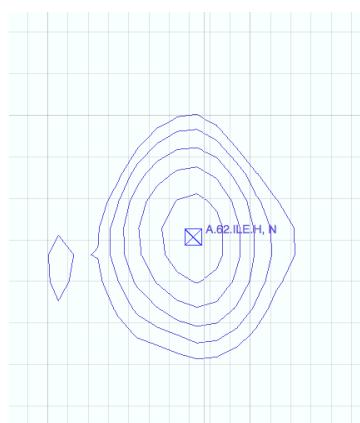
This peak list was copied from Tstar without ligand, therefore, some peaks will have changed position upon titration with the ligand. You will have to find the shifted peaks and correct their position.

3c Move peaks manually.

- Locate a single peak and zoom in using the middle mouse wheel, until it's in the middle of the display. Hold down the left mouse button and drag to pan the spectrum.
- On the peak table, double click on a row, e.g. # 10 | A.62.ILE. The spectrum display will navigate to the selected peak coordinates.
- Move the mouse cursor on top the selected peak [X]: hold down the **middle mouse wheel button or right** and drag across the selected peak. The peak will start to move, when you are in the correct position, release the mouse button.
- Use the shortcut "SE" to centre the peak to its extremum.

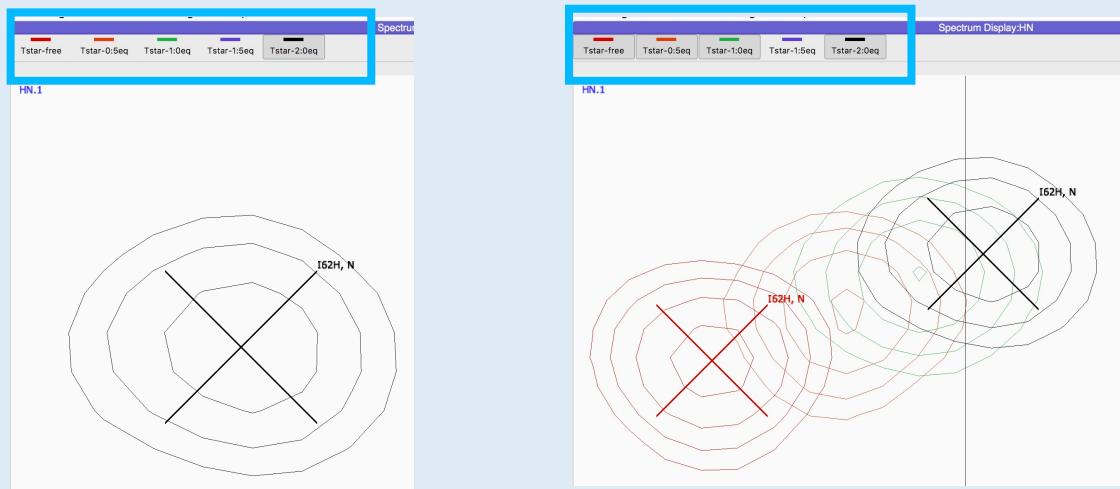


hold middle or right mouse
and drag



Release

Move Peaks



To change the order how spectra are displayed: Middle mouse on the toolbar button and drag and drop the button to a new positon in the toolbar

3D Map the trajectory.

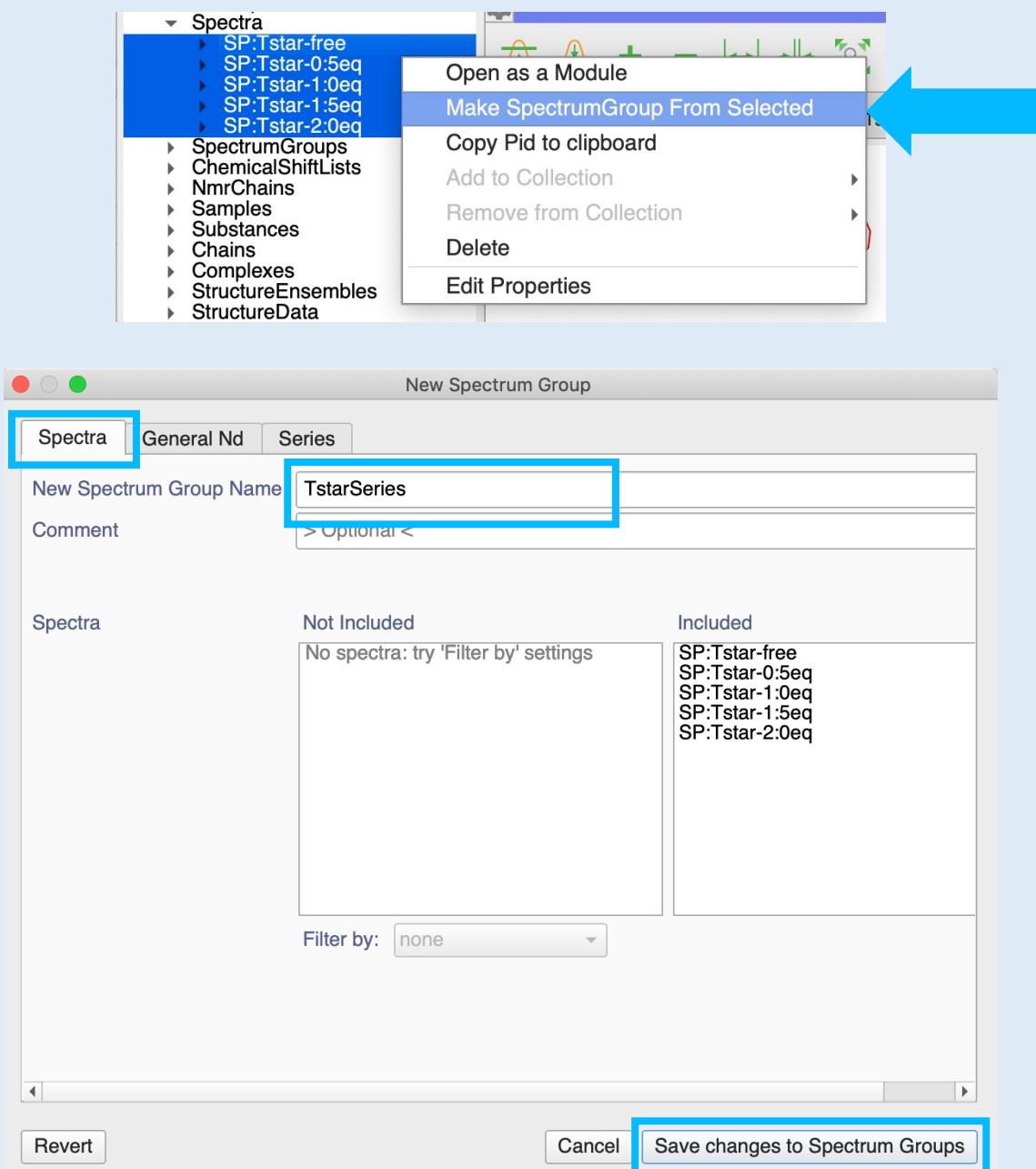
To map the trajectory of the peak shifts it is important to check the other spectra too.

- toggle on and off manually the spectra on the toolbar to see any differences in the peak positions across the different spectra.

Useful shortcuts:

- “Tab–Tab” (press twice →→)
- Displays the next spectrum and hides all others
- “Tab–A” : Displays all spectra
- “Tab–X” : Reverts displayed spectra

Group Spectra

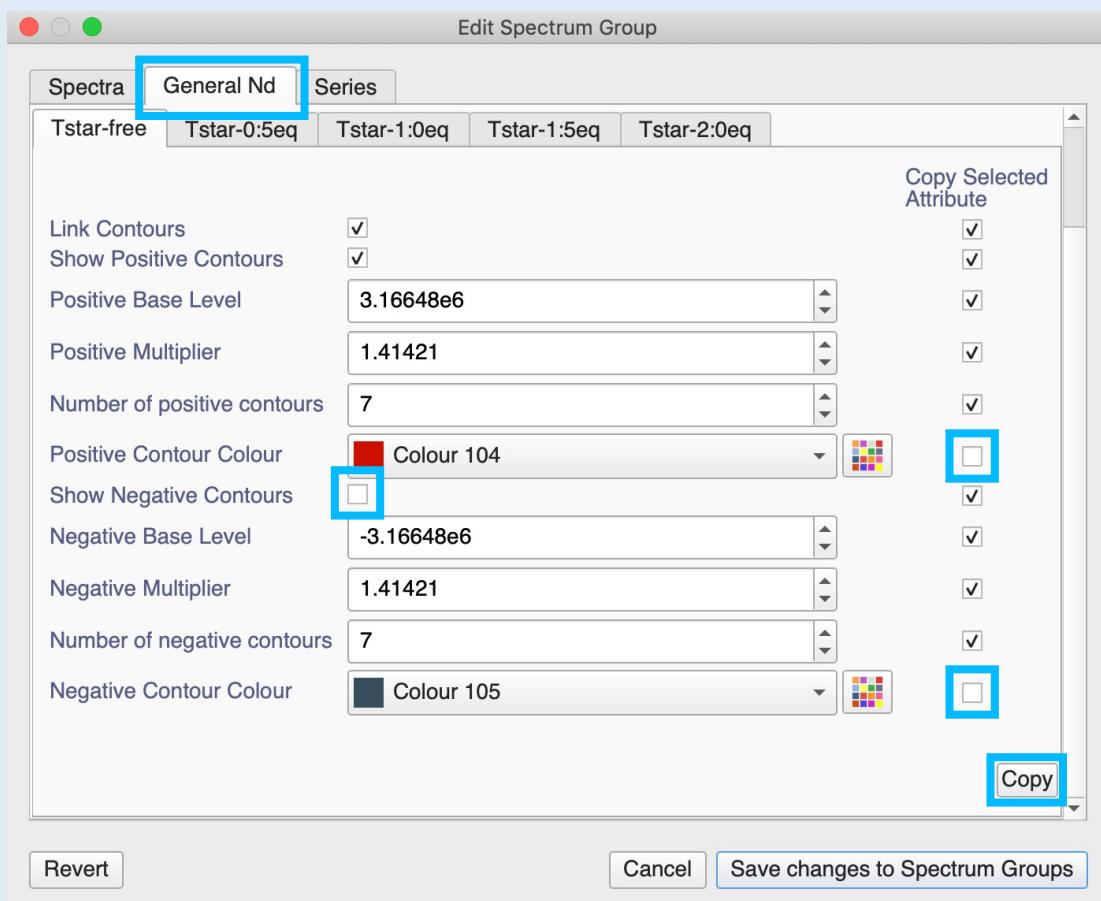


3E Create spectrum group.

For further analysis spectra have to be organised as a spectrum group.

- Select all spectra in the project.
- Right click on *Make SpectrumGroup From Selected*
- In the pop up window type the name of the Spectrum Group and click **Save changes to Spectrum Groups**.

Group Spectra



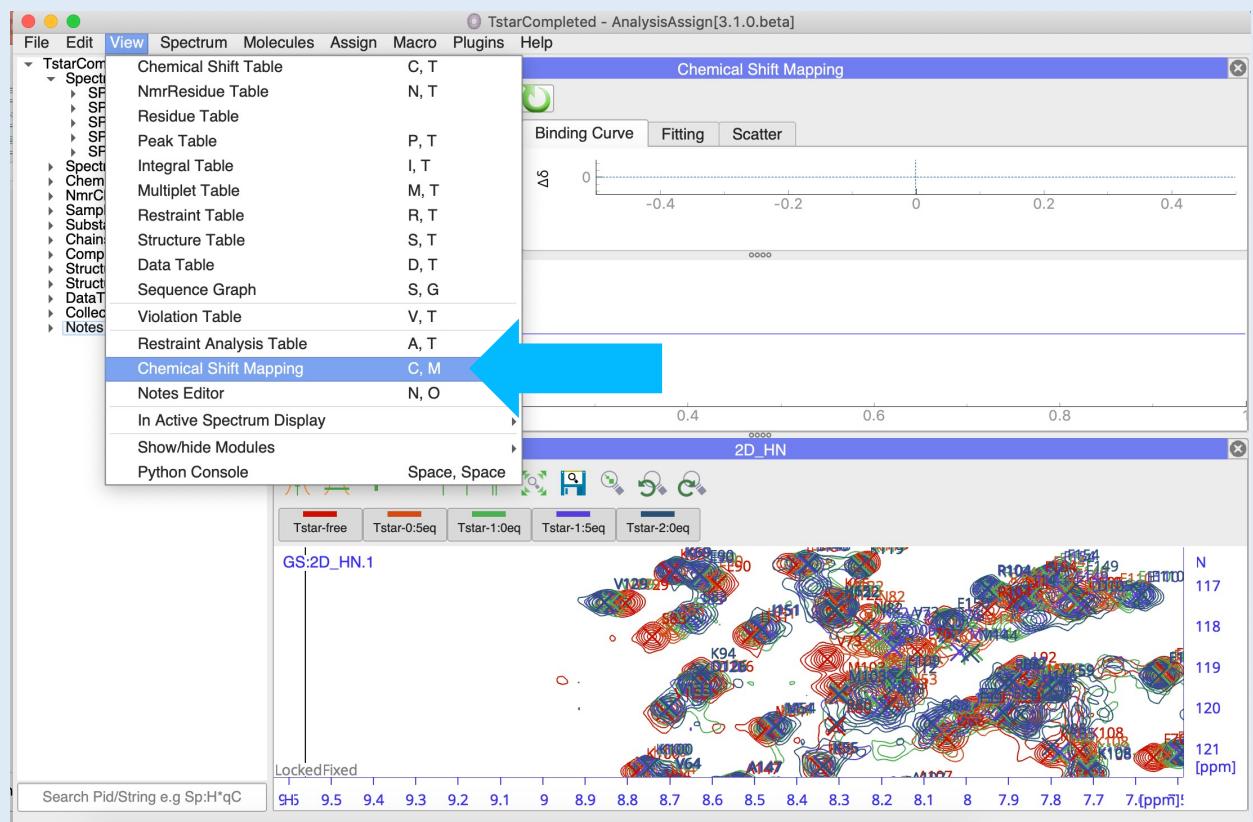
3F Change parameters across spectrum group.

You can copy parameters between the spectra using the pop up window:

- Click on the General ND tab
- Untick Positive Contour Colour, Show Negative Contours, Negative Contours Colour and **confirm by clicking *Copy*.**

To have a full overview of the ChemicalShiftMapping module, you will need to repeat points 3D and 3E for as many peaks as possible. The project “TstarCompleted ccpn” includes fitted peaks for all spectra and created spectrum group, you might continue the tutorial using that project instead.

Map Peaks



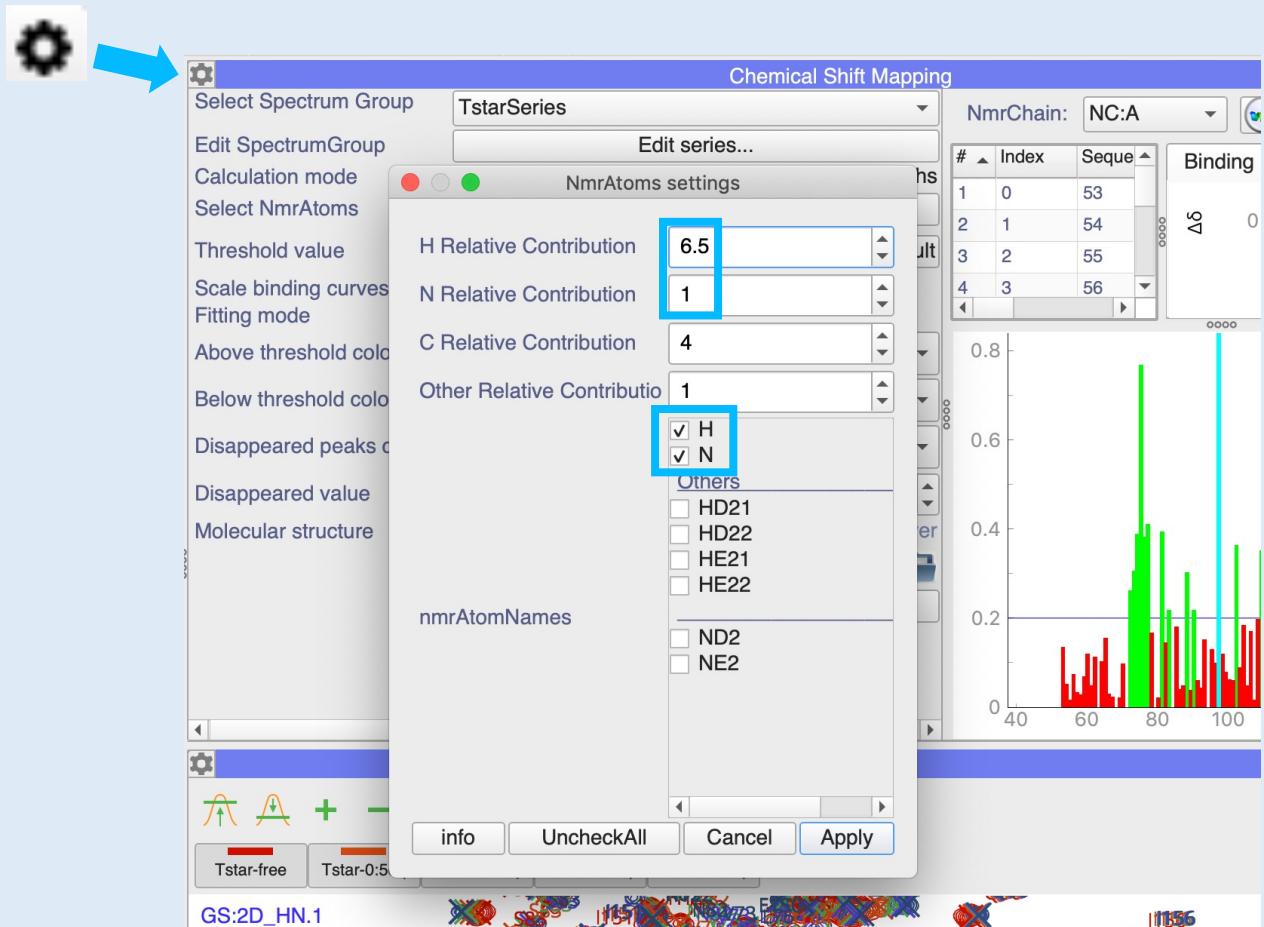
The **Chemical Shift Mapping module** includes a bar chart plot, a residue table and a settings panel. As previously stated, each peak represents a single residue in the protein (apart from the sidechain peaks). From now on, we will refer to a residue assigned to its spectrum peak as an **NmrResidue**. The NmrResidues that have moved above a threshold of 0.1 ppm have been plotted in green; the others in red.

4A Open the Chemical Shift Mapping module, shortcut “CM”.

Once you have moved all peaks correctly, inspect the chemical shift perturbations using the Chemical Shift Mapping module.

- Go to the main menu → *View* → *Chemical Shift Mapping* module or use the shortcut “CM”.

Map Peaks



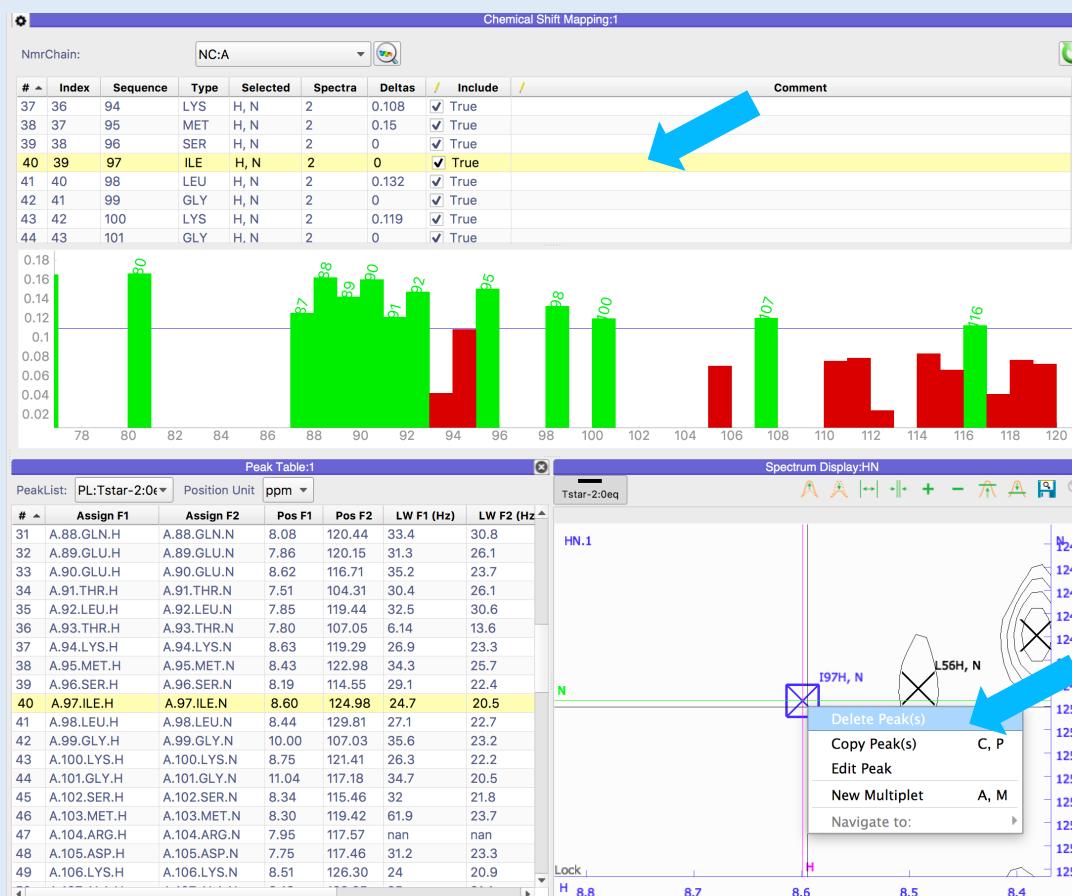
4B Changing the default values.

Click the “gear icon” in the top left corner of the module .

Set:

- Click *Select NmrAtoms -> Relative Contribution*: H = 6.5, N = 1 , names as default
- Calculation mode*: position
- Threshold line*: 0.200 ppm
- You might want to change the plot colours
- PDB file Path*: click the folder icon  and select TSTAR-KH.pdb in the CcpnCSP_tutorial/pdbStructure directory.
- Click the *Update All* button.**

Map Peaks (Optional)



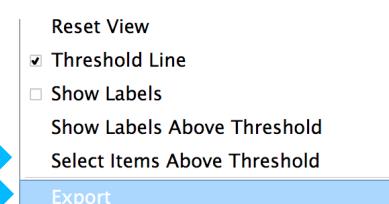
4C Find the disappeared Peak.

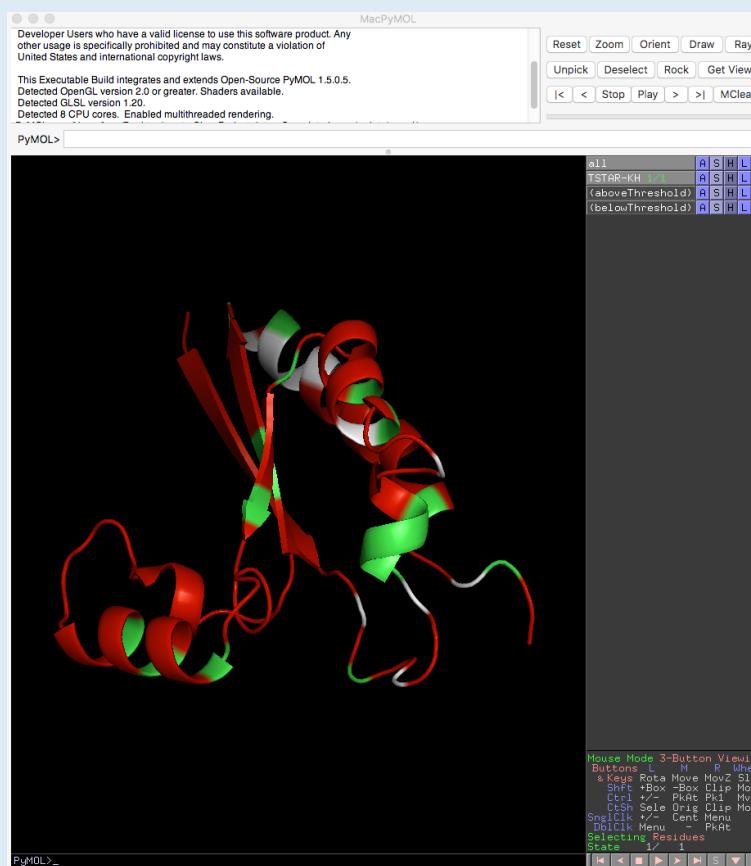
In this protein a peak has disappeared after the titration with the ligand.

- If present in the plot, double click on the residue bar 97 to navigate to the residue coordinates on the spectrum display, otherwise double click on the peak table #40 | A.97ILE.H | A.97ILE.N.
- Right click and Delete the peak from the spectrum Tstar-2:00eq You can add a comment on the *Comment* column Chemical Shift Mapping table cell by double clicking inside the cell.
- The plot will automatically refresh and mark the residue as "missing"

4D Export plot.

- Right click in the plot in an empty space
- Click *Show Label Above Threshold*
- Click **Export**
- On the Export popup Select Image File, then click export
- Save the file





PyMOL is a Python-enhanced molecular graphics tool. It excels at 3D visualisation of proteins, small molecules, density, surfaces. The visualisation through PyMOL of the perturbed residues on the surface's protein can be started using CcpNmr. Download and install from <https://pymolwiki.org/index.php/MAC>. Install

5A Link PyMol to CcpNmr Analysis

- Main menu → Project → Preferences...
 - click on External Programs tab
 - in PyMol, click on the folder icon,
 - search on your disk for the executable PyMol file, on a Mac this will be on:
/Applications/MacPyMOL.app/Contents/MacOS/MacPyMOL
 - click on Test, Pymol will start if has been linked successfully
 - click *OK*

5B Click “Show on Molecular Viewer”.

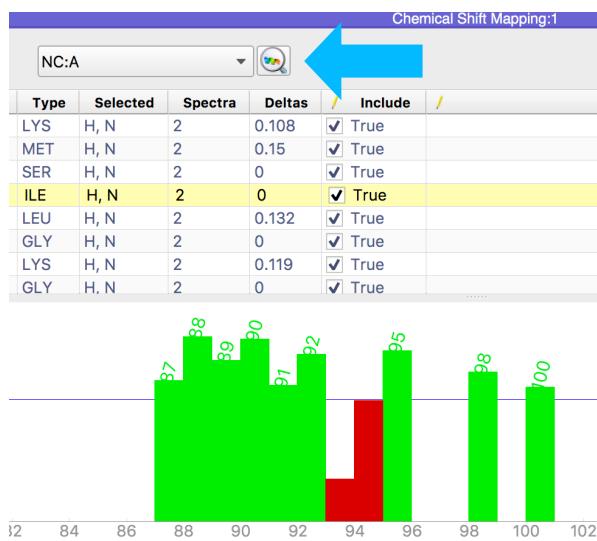
By clicking the button



on the Chemical Shift mapping inside CcpnNmr Analysis, PyMol will be launched with the Tstar structure already loaded in. In Pymol two new selections will be created:

- (*aboveThreshold*) , same colour as the bars in CcpNmr
- (*belowThreshold*) , same colour as the bars in CcpNmr
- (*missing*) , same colour as the bars in CcpNmr

You will be able now to analyse graphically the residues that are involved in the interaction with the ligand.

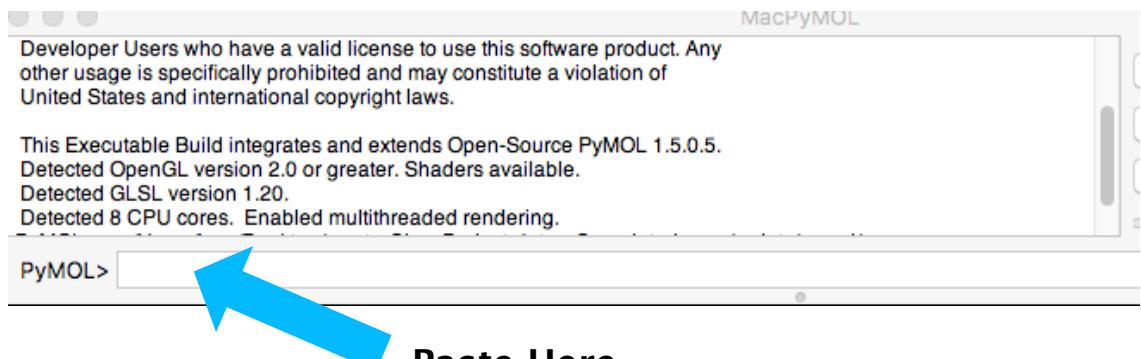


5C (Optional) Export image with transparent background

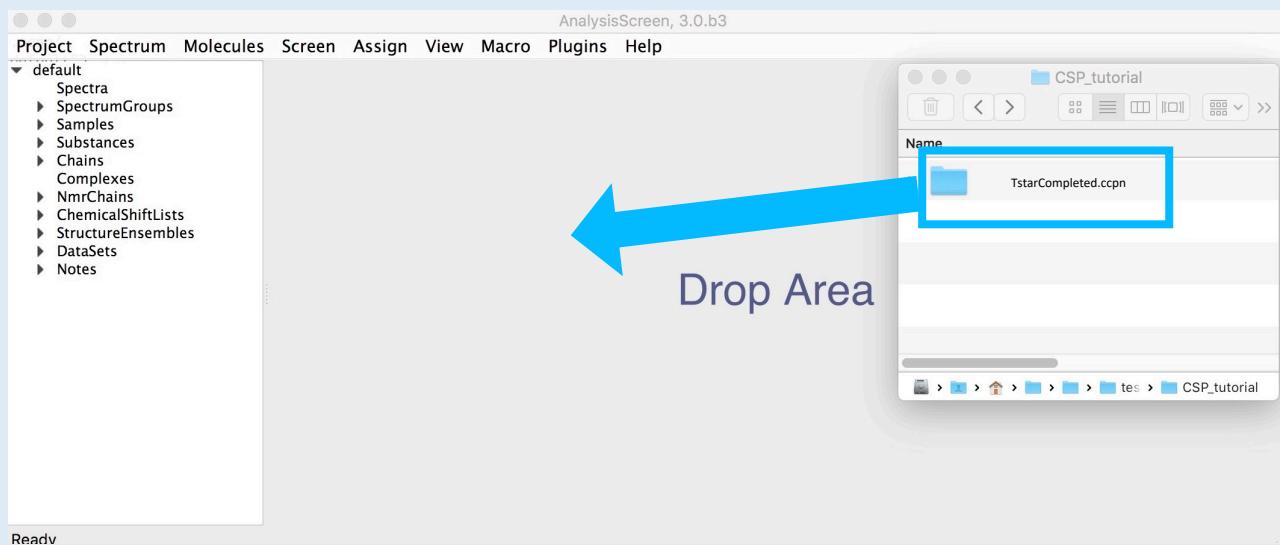
Copy and paste these commands on the Pymol terminal:

- `set ray_opaque_background, 0`
- `png ~/Desktop/tStar.png, width=1000, dpi=300, ray=1`
- press enter

Find the image on your desktop



Binding Curves



6A Drag & drop “TstarCompleted ccpn” (uncompressed) into the sidebar.

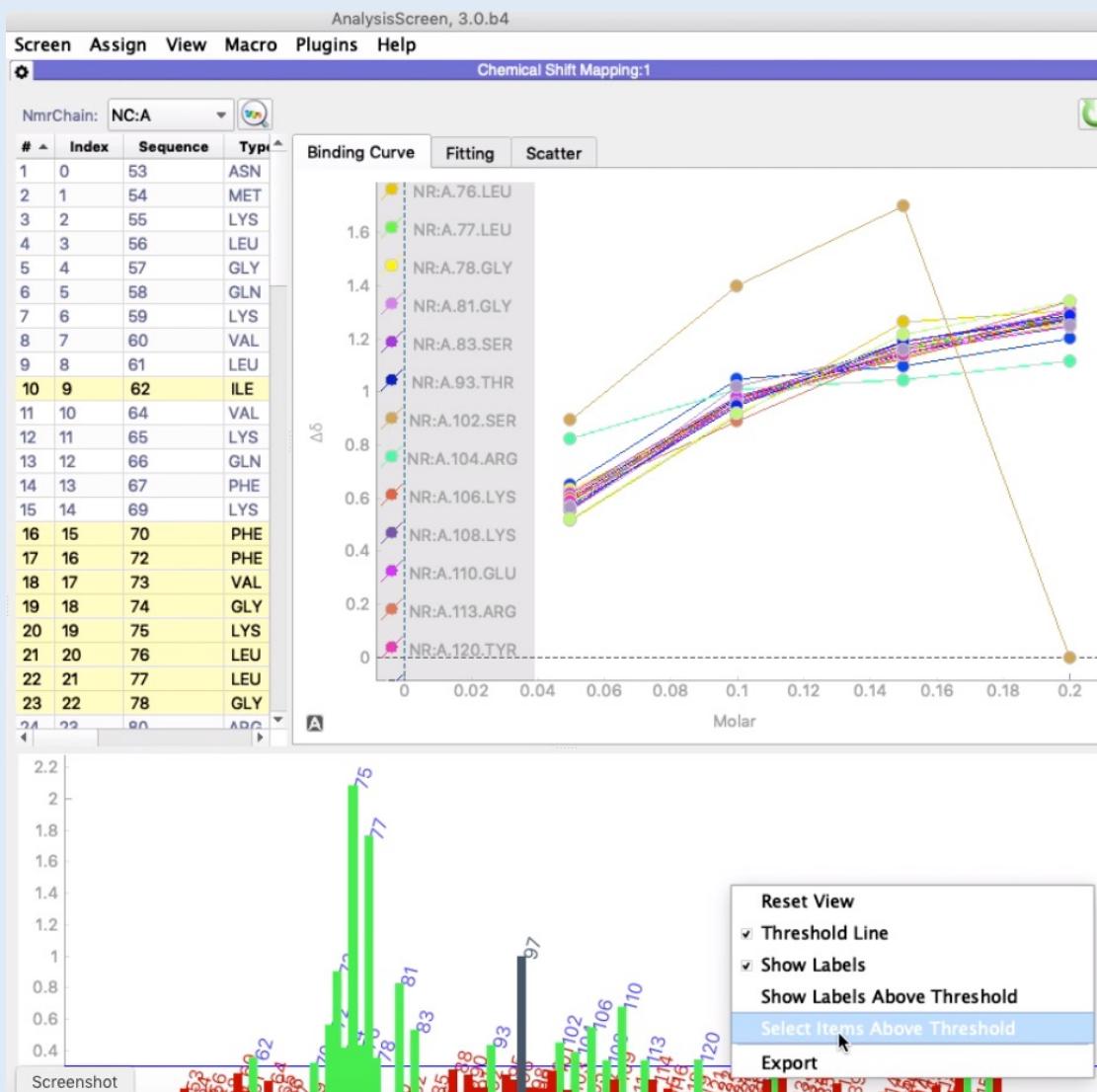
This project contains peak lists in all the spectra with already fitted peaks.

If not already opened, load the **TstarCompleted** project and press “CM” to display the Chemical shift mapping module.

Display all five spectra in a single spectrum display and zoom to focus on a single peak.

6

Binding Curves



The binding plot is created based on $\Delta\delta$ for each spectrum recorded at different ligand concentrations. This functionality is also useful to spot fitting errors as described below.

6B

Select NmrResidues above threshold

Right click on the bar graph canvas and click *Select above threshold*.

All the binding curves for the selected NmrResidues will be plotted.

You might see some of the curves don't follow a "one site binding" curve perfectly. It could be some of the peaks are not correctly fitted. E.g. the nmrResidue 102.SER

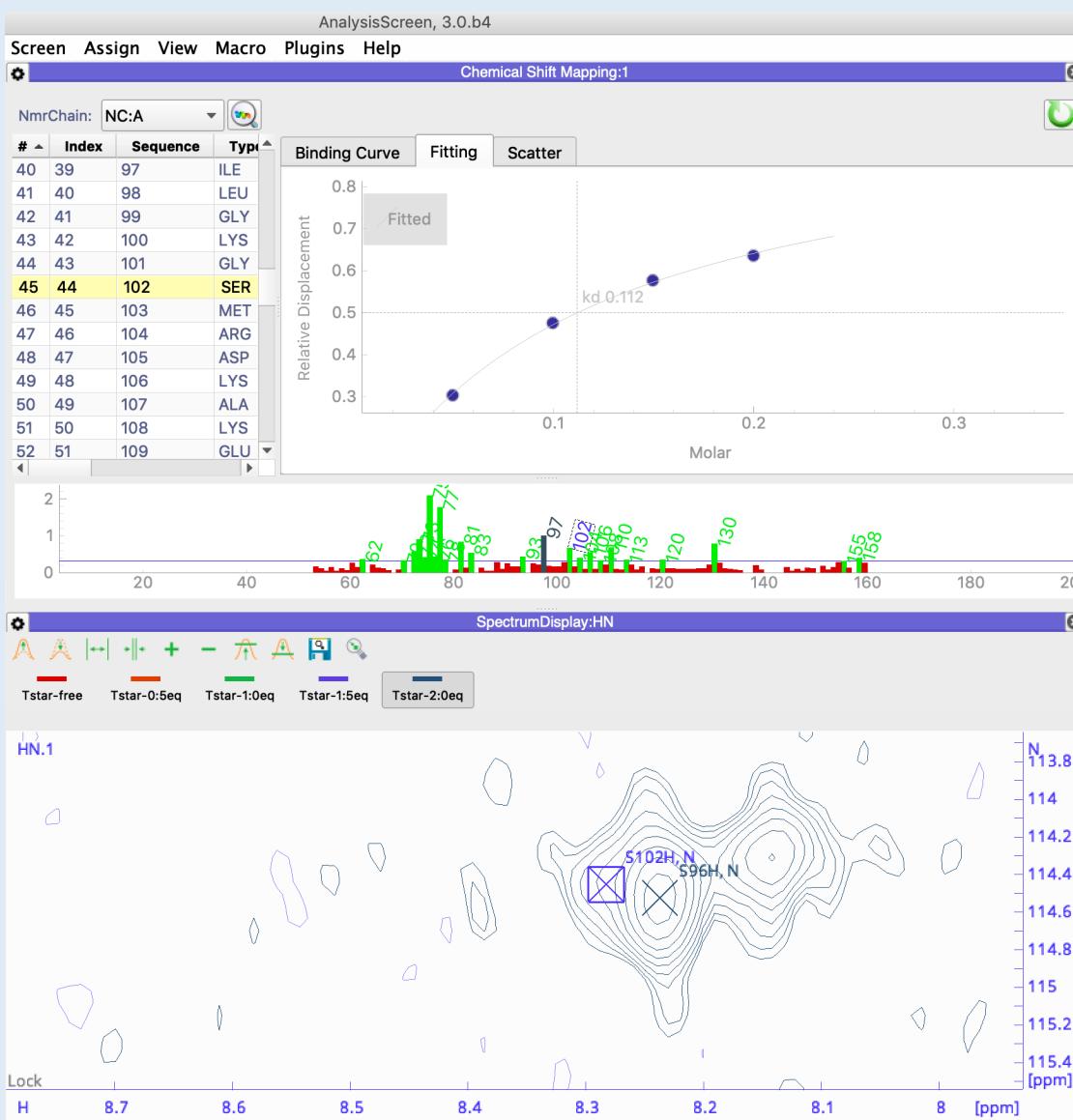
6C

Double click any of the point for the misplaced curve. (Colours may vary)

This will navigate to NmrResidue on the spectrumDisplay; using the "Tab-Tab" shortcut follow the peak.

6

Binding Curves



6C

Move the peak for the spectrum Tstar-2:0eq.

This peak has an ambiguous position, once you place it, click the "update" button

This will update the table and plots.

Click on the Fitting tab to inspect the estimated Kd for the select NmrResidue.

To exclude an NmrResidue from the multisection:

- Hold Ctrl (Cmd on a Mac) and click the item on the table.

1. Empty bar chart:

- Check the selected NmrChain contains valid NmrResidues and nmrAtoms are assigned to peaks in at least two different spectra PeakLists:
- SequenceCode for the NmrAtom is a numerical digit, e.g. sequenceCode=1
- Check the peaks in the selected spectra (settings) are fitted correctly and have differences (position, height etc.) across the different spectra.
- Check the selected NmrAtoms (settings) for ambiguous combinations. E.g. selecting H, H1, H2, Nh3 simultaneously might not give any results

1. Empty Binding curves:

- Check at least three or more spectra (containing fitted peaks) are selected on the settings panel
- Check Ddelta is not None or 0. See 1A2

1. Peaks not “snapping” to extrema (“SE” shortcut):

- Shortcut out-of-sync. Try any pressing another key, then re-do “SE”
- Peak is too far from the extremum. Move closer with the mouse or keyboard commands

2. PyMol not opening:

- Check PyMol has been correctly linked to Analysis as 5A.
- Alternatively, You can run the PyMol script directly from PyMol. The CcpNmr scripts are located inside the project folder. For PyMol will be on: name ccpn > script > pymol > chemicalShiftMapping_Pymol_Template.py Simply, run this file from the PyMol File menu.

Contact Us

Website:

www ccpn.ac.uk

Suggestions and comments:

support@ccpn.ac.uk

Issues and bug report:

<https://forum.ccpn.ac.uk/>

Cite Us

Simple High-Resolution NMR Spectroscopy as A Tool in Molecular Biology. LG Mureddu and GW Vuister. *FEBS Journal* (2019). [doi:10.1111/febs.14771](https://doi.org/10.1111/febs.14771)

CcpNmr AnalysisAssign: a flexible platform for integrated NMR analysis. SP Skinner *et al.* *J. Biomol. NMR* (2016). [doi:10.1007/s10858-016-0060-y](https://doi.org/10.1007/s10858-016-0060-y)