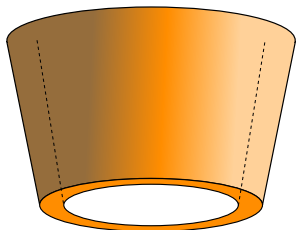
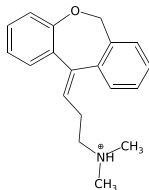
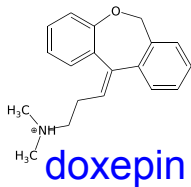


# Revealing ligand-receptor interaction NMR titration

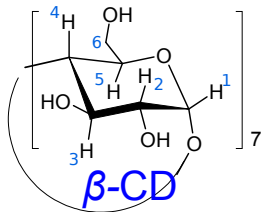
Jan Novotný

May 3, 2017

# Introduction of reacting partners

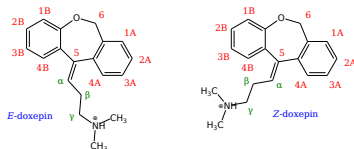


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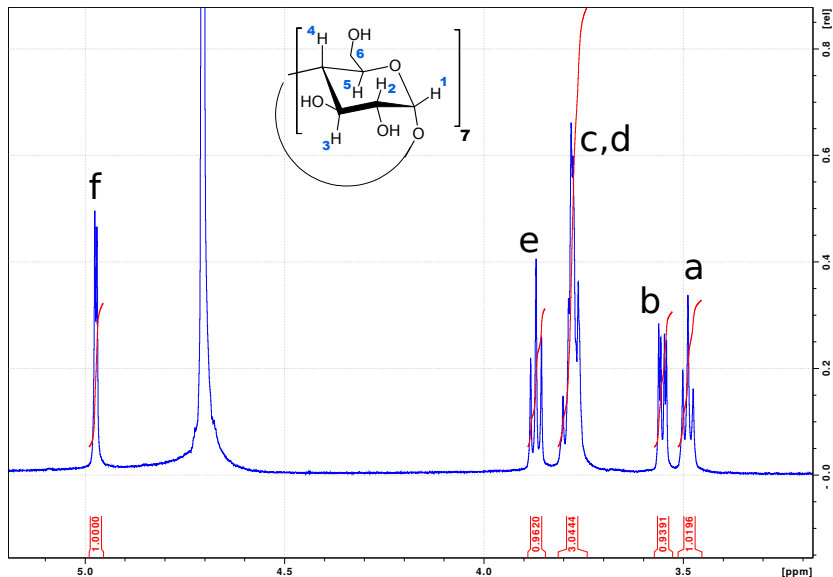


# Recommended procedure

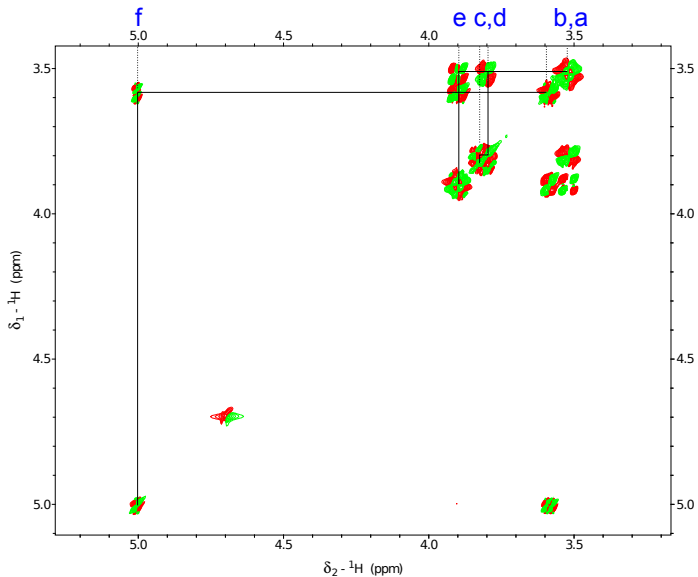
- 1 Assignment of free receptor:  $\beta$ -cyclodextrine (1D  $^1\text{H}$ , DQF-COSY)
- 2 Assignment of free ligand: doxepine (1D  $^1\text{H}$ , DQF-COSY, NOESY)
  - Identification of proton resonances of ring A and B
  - Determination of major and minor conformation of doxepine
- 3 1D NMR titration: rearrangement of  $\beta$ -cyclodextrin resonances upon interaction  $\Rightarrow$  identification of inner protons
- 4 1D NMR titration: rearrangement of doxepin resonances upon complexation  $\Rightarrow$  estimation of binding mode
- 5 ROESY spectrum of complex: ROE intermolecular contacts
- 6 Fitting the titration isotherm



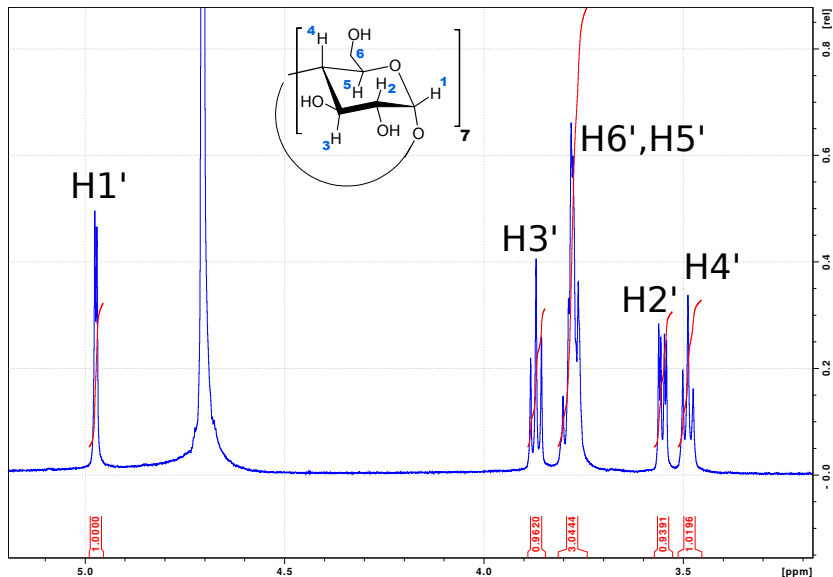
# 1D $^1\text{H}$ of $\beta$ -cyclodextrin in $\text{D}_2\text{O}$



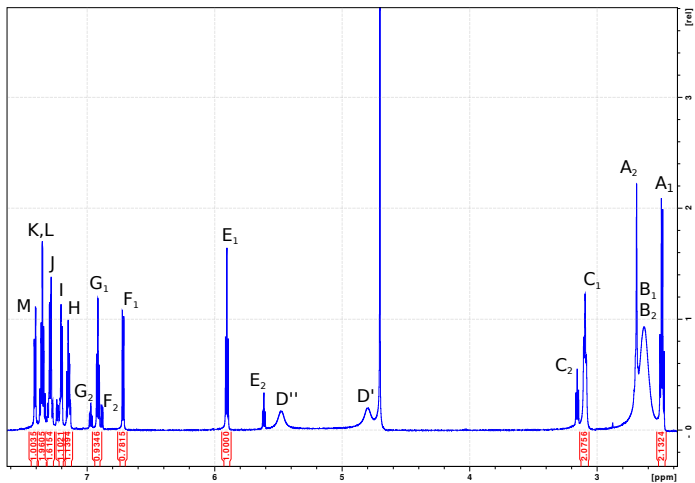
# DQF-COSY of $\beta$ -cyclodextrin in $D_2O$



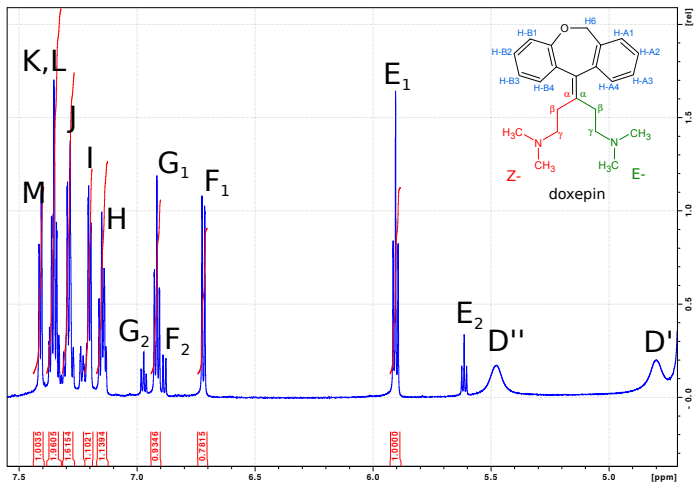
# 1D $^1\text{H}$ of $\beta$ -cyclodextrin in $\text{D}_2\text{O}$



# 1D $^1\text{H}$ of doxepin in $\text{D}_2\text{O}$

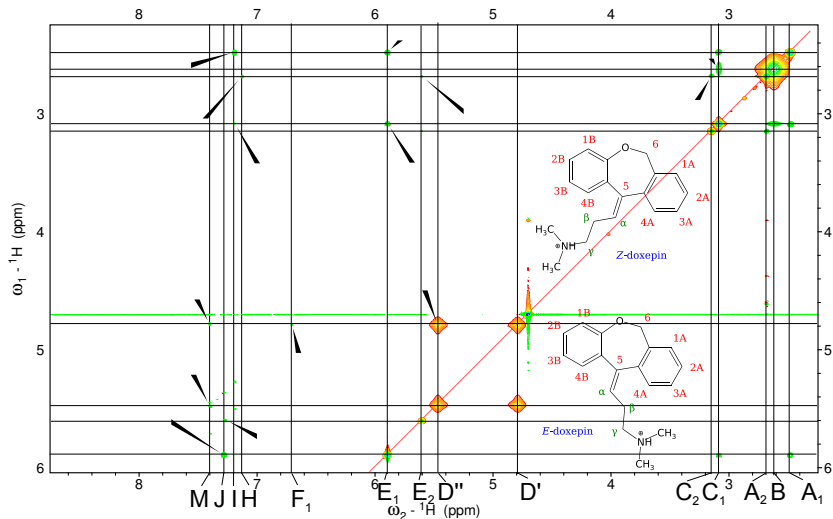


# 1D $^1\text{H}$ of doxepin in $\text{D}_2\text{O}$

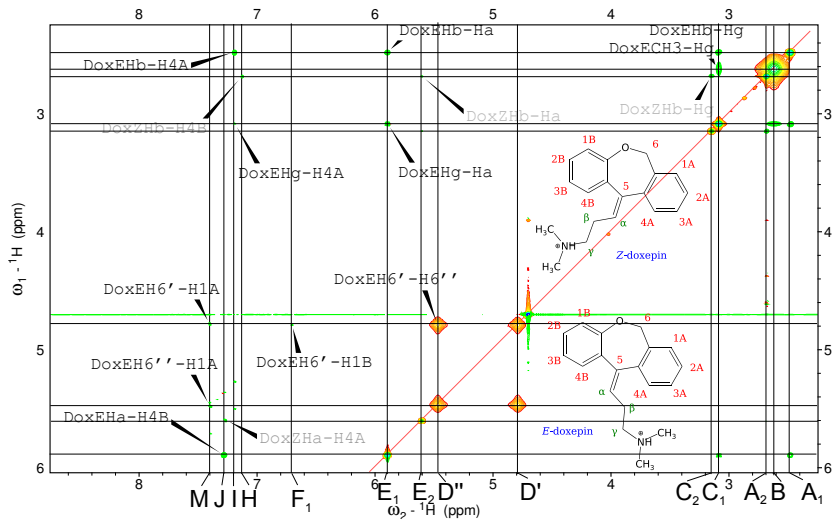




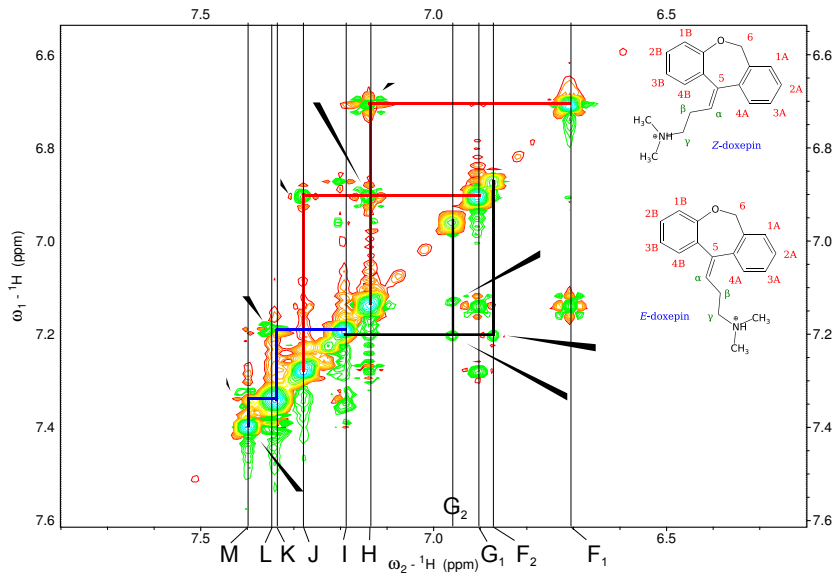
# NOESY 700ms of doxepin in D<sub>2</sub>O



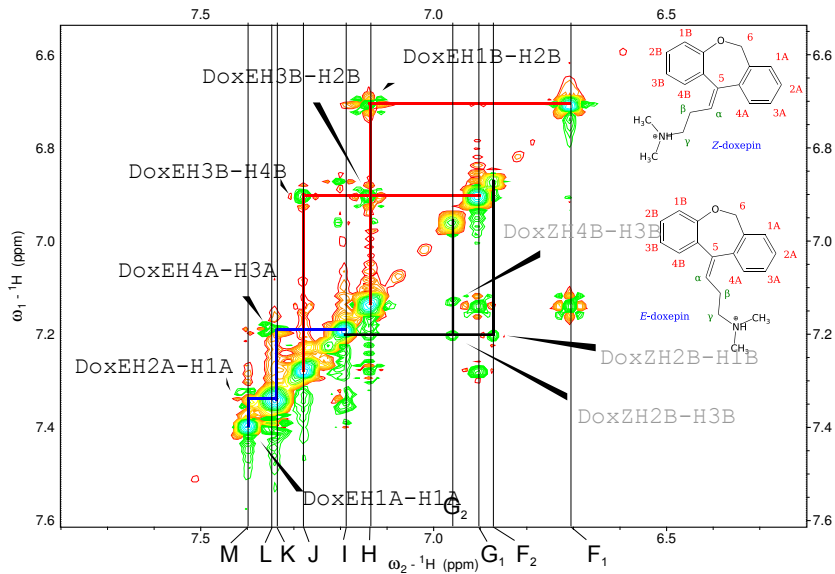
# NOESY 700ms of doxepin in D<sub>2</sub>O



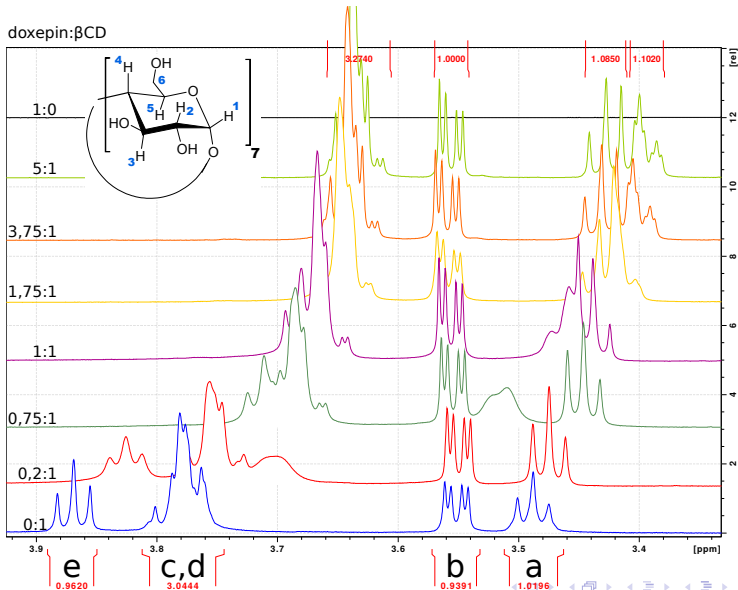
# NOESY 700ms of doxepin in D<sub>2</sub>O



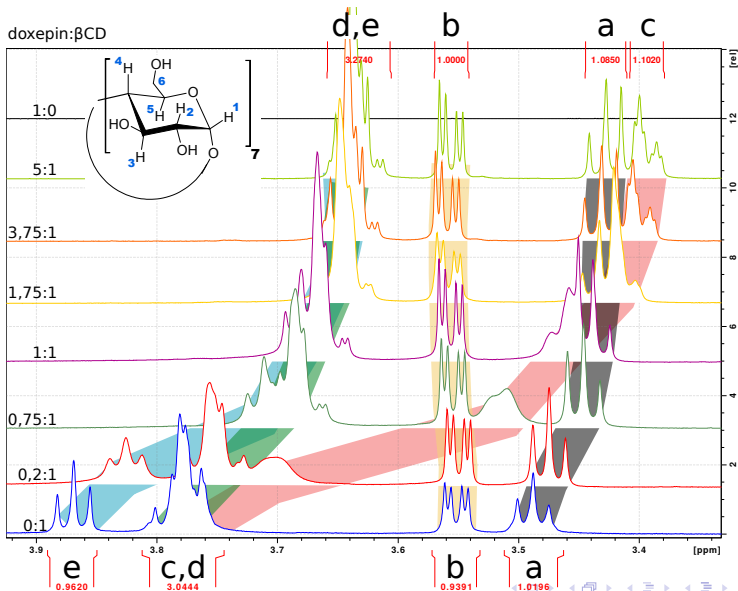
# NOESY 700ms of doxepin in D<sub>2</sub>O



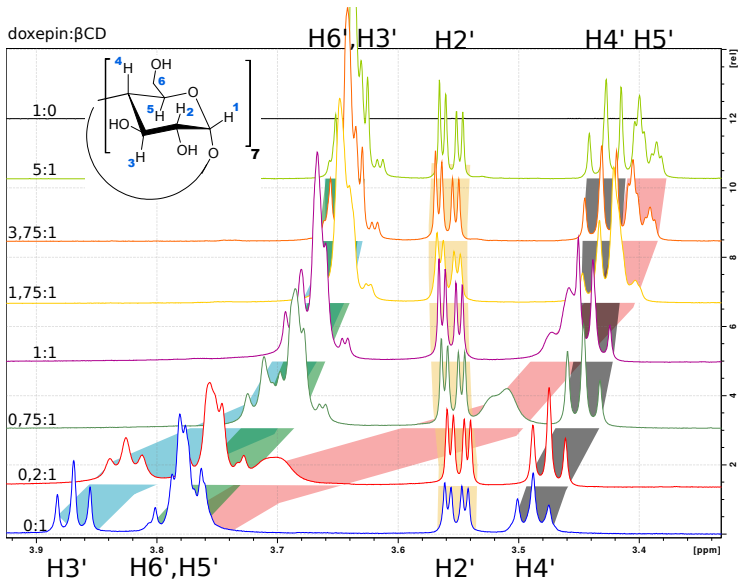
# 1D $^1\text{H}$ NMR titration: $\beta$ -cyclodextrin



# 1D $^1\text{H}$ NMR titration: $\beta$ -cyclodextrin

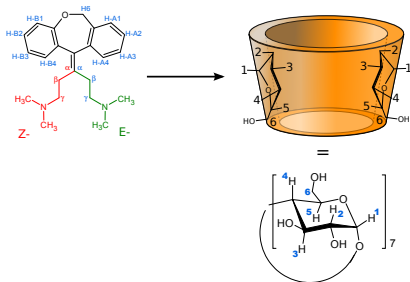
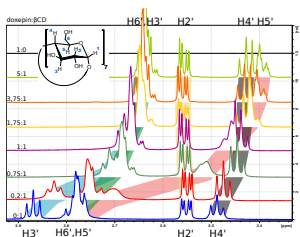


# 1D $^1\text{H}$ NMR titration: $\beta$ -cyclodextrin



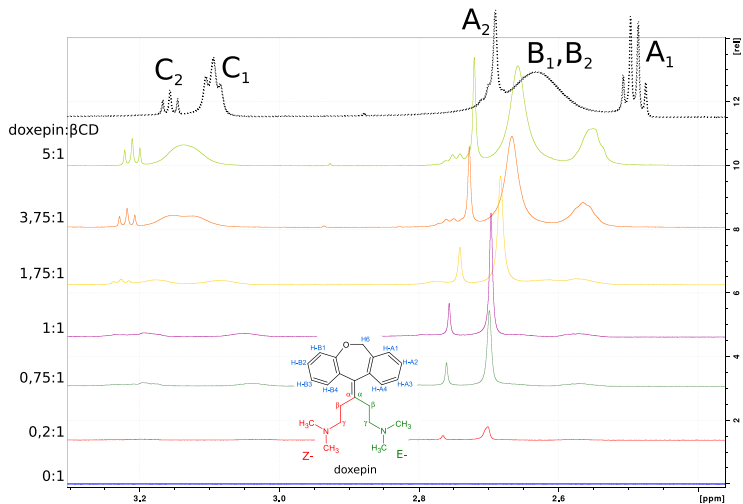
# 1D $^1\text{H}$ NMR titration: $\beta$ -cyclodextrin

Inner  $\beta$ -CD protons **H3'** and **H5'** are dominantly exposed to shielding induced by ring current of aromatic ring.

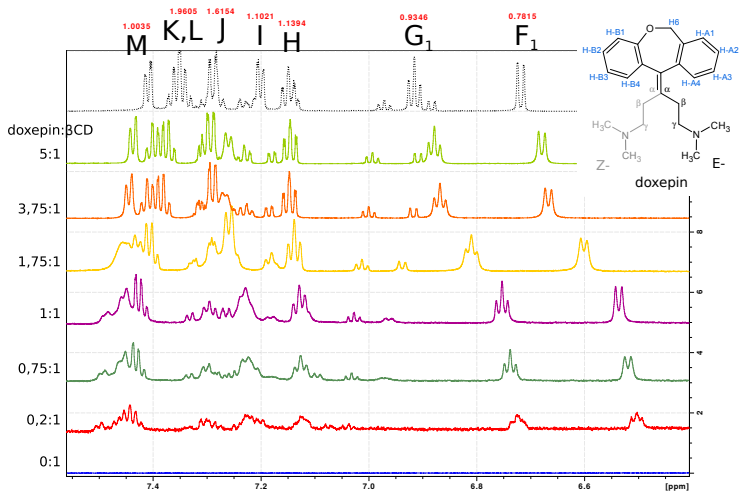




# 1D $^1\text{H}$ NMR titration: doxepin

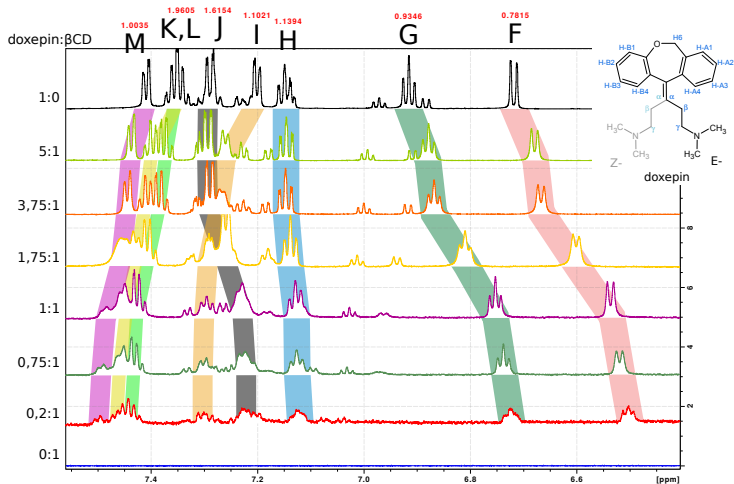


# 1D $^1\text{H}$ NMR titration: doxepin



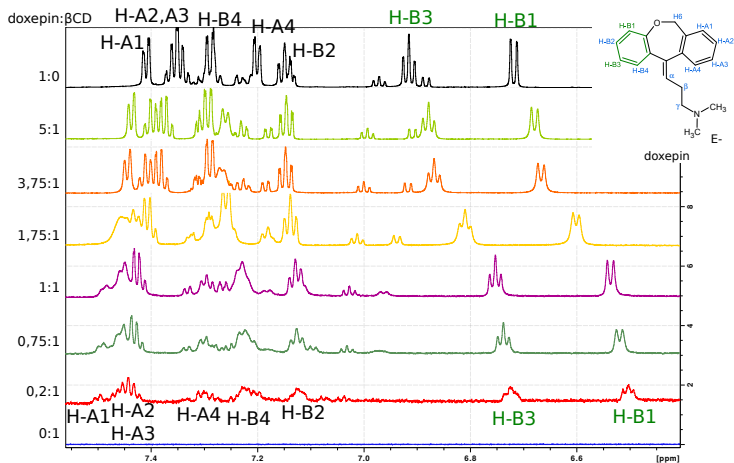
Find the most perturbed resonances and estimate the time regime of interaction:

# 1D $^1\text{H}$ NMR titration: doxepin



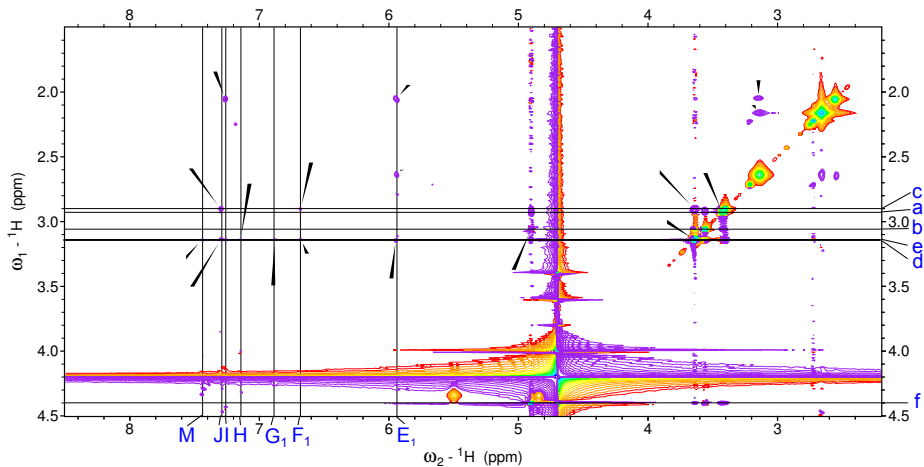
Find the most perturbed resonances and estimate the time regime of interaction:

# 1D $^1\text{H}$ NMR titration: doxepin

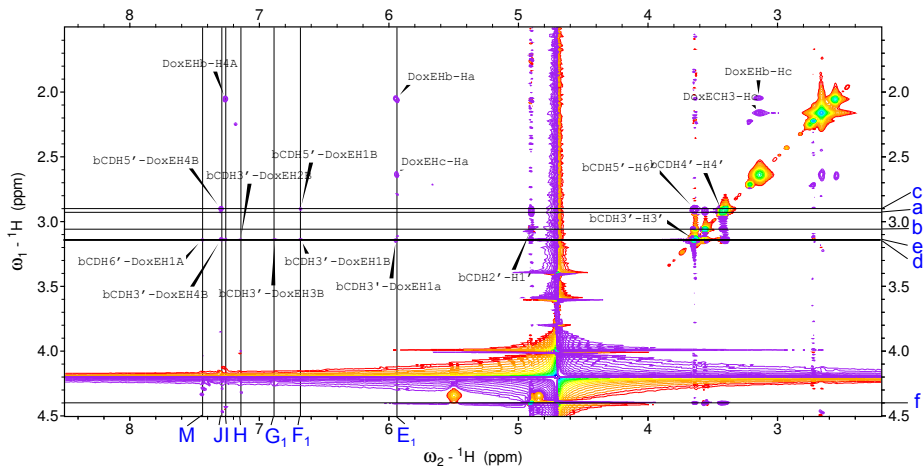


One set of broaden signals of doxepin  $\Rightarrow$  fast/intermediate rate of exchange.

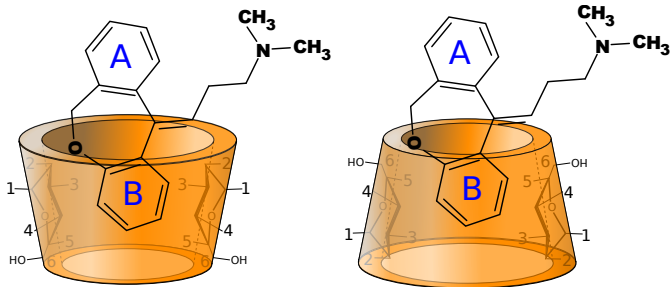
# ROESY 300ms - doxepin: $\beta$ -cyclodextrin=5:1



# ROESY 300ms - doxepin: $\beta$ -cyclodextrin=5:1

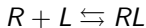


# Proposed orientation of doxepine in $\beta$ -cyclodextrin



# 1D $^1\text{H}$ NMR titration: determination of stoichiometry/ $K_a$

Titration curve of  $\Delta\delta(\text{H1}')$  as a function of increasing concentration of doxepine.



$$K_a = \frac{[RL]}{[R][L]} = \frac{[RL]}{(R_0 - [RL])(L_0 - [RL])}$$

Upon titration we are gradually changing  $L_0$  concentration:

$$\delta_R = [RL]\delta_{R_{bound}} + (R_0 - [RL])\delta_{R_{free}}$$

$$\Delta\delta_R = \delta_R - \delta_{R_{free}}$$

