Development of sampling method for cerebrospinal fluid in mice and effect of sampling method on drug concentration in cerebrospinal fluid

## 無断転載禁止

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## Purpose

Since the sampling of cerebrospinal fluid (CSF) from mice was difficult due to mouse body size, we developed a sampling method and a novel sampling device for the CSF from mice. We investigated the contamination ratio of blood to the CSF due to the sampling method. Furthermore, The effect of the contamination on drug concentration in the CSF was also evaluated by using tool compounds.

#### **Materials and Methods**

<u>Development of sampling method for cerebrospinal fluid in mice:</u> Mice and rats were anesthetized by inhalation of isoflurane, and the blood and the CSF were collected from the caudal vena cava and the cisterna magna, respectively. The weight of the collected CSF was measured. The CSF volume was estimated from the weight on the assumption of specific gravity as 1. Red blood cells in the blood and the CSF were counted. The contamination ratio of the CSF by the blood was calculated as the number ratio of red blood cells in the CSF (RBC  $_{\rm CSF}$ ) to those in the blood (RBC  $_{\rm blood}$ ).

Effect of blood contamination on drug concentration in CSF (Ccsf): Metformin, ranitidine, ibuprofen, imipramine, glimepiride, glibenclamide and nefazodone were administered to mice (8W) intraperitoneally. Plasma, brain and CSF samples were obtained after dosing. CSF was collected deliberately contaminated with blood.

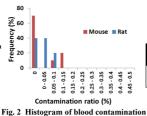
All animal protocols were approved by the Institutional Animal Care and Use Committee of Shonan Research Center, Takeda Pharmaceutical Company Ltd.

<u>Bioanalysis</u>: Concentrations of metformin, ranitidine, ibuprofen, imipramine, glimepiride, glibenclamide and nefazodone in samples were quantified by LC/MS/MS analysis.

<u>Unbound fraction:</u> Unbound fraction of compounds in plasma and brain was measured by equilibrium dialysis method (fu,p and fu,b, respectively)

### Results The novel method of mouse CSF collection





 Amount of CSF (μL)

 Mouse
 Rat

 Mean
 11.8
 159.2

 SD
 2.9
 26.2

Table 1 Collection amount of mouse

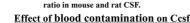
CSF and rat CSF

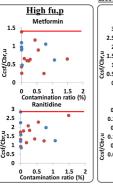
Contamination ratio =  $\frac{RB}{RBG}$ 

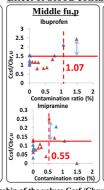
Table 2 Unbound fraction of 7 compounds in mouse plasma and brain

Compounds	fu,p	fu,b
Metformin	1.0	0.808
Ranitidine	1.0	0.571
Ibuprofe n	0.12	0.263
Imipramine	0.10	0.010
Glimepiride	0.030	0.077
Glibe nclamide	0.030	0.055
Nefazodone	0.020	0.004

from mice.







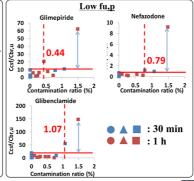


Fig. 4 Correlation between fu,p and increasing ratio by contamination in Cesf.

Max increasing ratio in Cesf was calculated by the following formula, Max increasing ratio in Cesf = [Cesf/Cbr,u]max/[Cesf/Cbr,u]nc. [Cesf/Cbr,u]max is maximum value of Cesf/Cbr,u. [Cesf/Cbr,u]ne is maximum value of Cesf/Cbr,u in samples without contamination.

# Fig. 3. Relationship of the values Ccsf/Cbr,u with contamination ratio (%). Cbr,u = Brain cone. x fu,b. Red lines indicate maximum Ccsf/Cbr,u at which contamination is 0%. Red dotted lines indicate minimum contamination ratio at which Ccsf/Cbr,u exceeds red line.

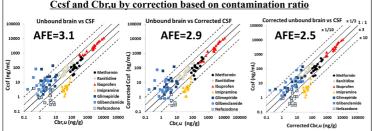


Fig. 5. The correlation between Ccsf and Cbr,u by correction based on contamination ratio Average-fold error (AFE) =  $10^{\circ}[1/N \times \Sigma[[\log (Predicted/Actual))]]$ 

# Corrected Cosf = CSF - Plasma conc. X Contamination ratio 1 - Contamination ratio Corrected Cbr,u = Cbr, u + Plasma conc. X Contamination ratio 1 + Contamination ratio

### **Summary**

- ✓ We established a method of CSF sampling from mice with maximum contamination ratio of 0.14%
- Minimum contamination ratio which affected compound concentration in CSF was 0.44%.
- The compounds with lower fu,p tended to show the larger effect on concentration ratio.
- ✓ AFE value between Ccsf and Cbr,u was improved by correction based on contamination ratio.

### Conclusion

We established a method of CSF sampling from mice applicable to the evaluation of compound concentration in CSF.

## COI disclosure information

We have no financial relationship to disclose for our presentation contents.