

## Fiche de proposition de stage de L3 et M1 UFR Chimie et Biologie

Si possible limitez-vous à 1 page (recto)

Une spécialité en M1: Master 1 Chimie du vivant. Pour les licences de chimie, deux mentions : C : Chimie, CB : Chimie et Biologie.

Cochez la spécialité correspondant à ce stage:

M1C

L3C

L3CB

Adresse et appartenance du laboratoire :

Institut de Biologie Structurale  
71, avenue des Martyrs, CS 10090 38044 Grenoble

Thématique générale du laboratoire ou du groupe de recherche (par mots clés)

Electron Microscopy and Method group directed by Dr. Guy Schoehn  
Research area: electron microscopy, negative strand RNA virus, hantavirus, nucleocapsid, RNA polymerase, replication, transcription.

Thème du stage proposé (en 10 lignes, si possible)

**TITRE : Structural characterization of Hantaan virus nucleocapsid by single particle cryo-electron microscopy and/or cellular electron microscopy**

**DESCRIPTION :**

Hantaviruses belong to the Bunyaviridae family, a large family of viruses that possess a segmented RNA genome of negative sense polarity. Each genome segment is enwrapped with the viral nucleoprotein N forming a ribonucleoprotein complex, called the nucleocapsid. This complex is essential in the viral cycle and protects the viral genetic information from the environment while providing a flexible helical template for viral transcription and replication by the viral RNA polymerase. The proposed Master I project will consist in the structural analysis of Hantaan virus nucleocapsids. Depending on his/her preference and on advances of the project at the time of the internship, the intern will be given two options:

1. express suitable constructs of recombinant Hantaan nucleoproteins, optimize their purification and analyse the resultant recombinant nucleocapsids by single-particle cryo-electron microscopy. We already have results showing that recombinant Hantaan nucleocapsids are straight helices amenable to high-resolution by cryo-EM.
2. Analyse fixed and inactivated Tula hantaviruses (non-pathogenic to humans) by cryo-electron tomography or analyse cells infected by Tula hantaviruses using cellular EM. Inactivated Tula hantaviruses and fixed infected cells will be provided by a collaborator.

Méthodologies et/ou techniques qui seront utilisées

Option1: biochemistry (protein expression, purification), negative stain and cryo-electron microscopy (initiation), image processing.

Option2: cryo-electron tomography, sub-tomogram averaging, cellular electron microscopy.

Personne à contacter (préciser si nécessaire les créneaux horaires) :

Name: Hélène Malet

Phone: 06 82 50 05 13

E-mail: [Complément d'information \(si nécessaire\) helene.malet@ibs.fr](mailto:Complément d'information (si nécessaire) helene.malet@ibs.fr)

Fiche de renseignement à retourner (en version pdf) par e-mail à :

[Olivier.Jarjayes@univ-grenoble-alpes.fr](mailto:Olivier.Jarjayes@univ-grenoble-alpes.fr) sous la forme

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