

**Offer type:** job

**Contract:** Temporary

**Salary range:**  $\geq 35,000$  and  $< 45,000\text{€}$  annual gross

**Employer:** Commissariat à l'Energie Atomique et aux Energies Alternatives

**Workplace:** Saclay - FRANCE

**Skill area:** Biology, Medicine, Health - **Biomolecules, pharmacology, therapeutics** - Chemistry

**Application deadline:** 03/01/2013. (extended)

SIMOPRO, Section of Molecular Protein Engineering, headquartered in Gif/Yvette nearby Paris, is a CEA leading research-focused Section headed by Dr. V. Dive in the field of protein/peptides engineering for diagnosis and therapeutic applications. SIMOPRO belongs to LERMIT laboratory of excellence on therapeutic innovation ([www.labex-lermit.fr](http://www.labex-lermit.fr)).

**Mission:**

POSTDOCTORAL POSITION in SIMOPRO/CEA:

Design and synthesis of MMP-12-specific probes as novel tracers for imaging of vascular inflammation in atherosclerosis.

**Who we are:**

The team of Vincent Dive has a long-standing expertise on the development of zinc-metalloproteinase inhibitors and probes through the exploitation of the phosphinic peptide chemistry. Our group has identified highly potent inhibitors able to selectively interact with MMP-12 (see Devel *et al.* 2006, 2010 and 2012). MMP-12 is a zinc Matrix MetalloProtease also called macrophage elastase that is secreted by macrophage and over expressed in various pathologies such as atherosclerosis. We recently demonstrated in a mice model of atherosclerosis with established plaques, that RXP470.1, one of our highly selective inhibitor, fully blocked further plaque growth and prevent plaque rupture (Johnson et al. ATVB 2011).

**The position:**

In the frame of the National Institutes of Health (NIH) funding, a post-doctoral position is available (24 months, open January 2013) for developing specific matrix metalloproteinase 12 probes. Based on the structure of highly selective MMP-12 inhibitors such as RP470.1 mentioned above, our aim is to design

novel MMP-12-specific tracers for imaging vessel wall inflammation in atherosclerosis. Bioavailability and targeting properties of such tracers will be evaluated by fluorescent imaging in different animal models where MMP-12 is over expressed.

**Candidates profile:**

As a PhD or a post-doctoral fellow you have expertise in organic chemistry and solid phase synthesis. You are familiar with basic biochemistry approaches to perform *in vitro* evaluation of tracers on a set MMPs and other Metalloproteases. You will be also taken to participate in *in vivo* fluorescent imaging experiments. You have a good command of spoken and written English. Age < 30 years old.

Applications will be sent to laurent.devel@cea.fr (email subject: in vivo fluorescence imaging) and will comprise a CV, an abstract of scientific researches -max. 2 pages- and 2 letters of recommendation. Only fulfilled application will be considered.