Newsflash, April 2016

Compliance with Good Laboratory Practices

As part of its continuing efforts to upgrade the quality services to clients and partners while complying with demand of regulatory authorities, **GenoSafe** has renewed once more its certificate for Good Laboratory Practices (GLP), receiving **status A GLP compliance**.

The GLP-laboratories, also called test facilities, conduct non-clinical safety studies. They are monitored on a two year cycle by the ANSM (the French National Agency for Medicines and Health Products Safety). The GLP Compliance Program includes, test facility inspections, study audits, follow-up inspections, and re-inspections. During the test facility Inspection not only the organization of the test facility but also on-going and completed studies are verified.

GenoSafe, a GLP-Laboratory since 2008, was inspected mid-February this year and is re-approved on April 2016 with Status A.
GenoSafe compliance relates to the testing of medicinal products.

About GenoSafe:
GenoSafe is a full contract research and consulting organization which specializes in evaluating the quality, safety and efficacy of gene and cell therapy products.
We assist our clients with customized services all along the development of their product, from the proof of concept (POC) through to market launch.
Our activities encompass 4 main domains of activities:

- Preclinical development
- Quality Control testing – Product characterization
- Clinical Development – Follow-up of subjects
- Regulatory Affairs consulting