**To**: GMO contact list

**Subject**: Consultation for a medicinal product for human use containing or consisting of a GMO - [name of product (INN)]

**Message:**

Dear colleagues,

In relation to the consultation hold with the Competent Authorities established under Council Directive 2001/18/EC with respect to the evaluation of environmental aspects, EMA inform you that European Public Assessment Report (EPAR) has been published: <link to the EPAR on the EMA Internet site>.

The EPAR reflects the scientific basis for the conclusion reached by the [Committee for Advanced Therapies (CAT) and] Committee for Medicinal Products for Human Use (CHMP) at the end of the centralised evaluation process, the approved Product Information in all EU languages as well as a the summary of the Risk Management Plan.

Yours sincerely,

[Add NAME]

Product Lead

European Medicines Agency