Lithuanian States Medicines Control Agency (human regulatory authority) to the Public Consultation Paper on the review of Regulation (EC) 1234/2008 to the handling of variations to purely national marketing authorisations.

1.	YES
2.	B
3.	YES
4.	Variations concerning changes to the active substance for the purposes of the annual update of a human influenza vaccine.
5.	YES
6.	YES
7.	YES
8.	YES, because workload is also increased.
9.	NO