



MINISTRY OF HEALTH  
OF THE CZECH REPUBLIC

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Prague, 13.10.2016  
Our ref.: MZDR 57844/2016/SOZ



**MZDRX00WD9AQ**

### **Biocide product notification**

Regarding your biocidal product notification for

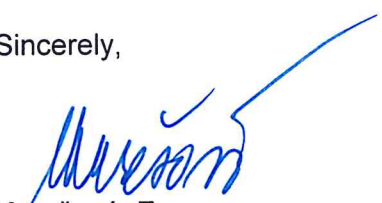
### **CIPEX 10 E**

as stipulated by Article 35 of Act No 120/2002 Coll., on Conditions for Placing Biocidal Products and Active Substances on the Market and on the Amendment of Some Related Acts, as amended, of 29.9.2016, please be advised that you have notified all data required pursuant to Article 35 of the above mentioned Act for biocide product notification.

Furthermore please be advised that Safety Data Sheets, proposed labelling, IFUs and efficacy tests attached to the notification in compliance with Article 35 of the above mentioned Act are not assessed or checked by the Ministry of Health, as the responsibility for the accuracy and completeness of data contained therein lies fully with the manufacturer or importer, or, where applicable, the distributor who places the biocide product on the domestic market.

Other responsibilities established by the quoted Act and by other legal regulations are not hereby prejudiced.

Yours Sincerely,



Pavla Marešová, Eng.  
on behalf of Director of Department of Strategy  
and Management of Public Health



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