MINISTRY OF HEALTH
OF THE CZECH REPUBLIC

BLEU LINE S.r.l.
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Prague, 13.10.2016
Our ref.: MZDR 57846/2016/SOZ

MZDRX00WD9DB

Biocide product notification

Regarding your biocidal product notification for

DEADYNA

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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