January 2014

BPR Compliance

To whom it may concern,

Addmaster, suppliers of the Biomaster range of additives can confirm that the active ingredients incorporated are supported for review under the requirements of the Biocidal Product Regulation – EU No. 528/2012.

The active ingredients are still going through the review process; until this is completed no biocidal products can be registered.

The current requirement for customers is to label any Treated Articles placed onto the market after the 1st September 2013.

These requirements include the following:

- All treated articles will need to be labelled, the labelling needs to be clearly visible, legible and durable.
- The labelling can be printed on packaging, instruction for use or warranty if the size or function of the article deems it necessary.
- The labelling does need to be in the language of the country of sale.
- The labelling requirements only apply to the company placing the treated article onto the market for the first time. There are no further obligations down the supply chain for the label to remain on the products for its life cycle

Any questions can be directed to the contact details below

Signed for and on behalf of Addmaster (UK) Ltd

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