

January 30th, 2015

BGMA Response to Branded Medicines Consultation

In reply to the Government's response to the results of the Consultation on Amendments to the Statutory Scheme to Control the Prices of Branded Health Service Medicines, Warwick Smith, Director General of the British Generic Manufacturers' Association (BGMA), said: "We welcome the Government making such a detailed response and are heartened that the Department of Health will not introduce an additional price cut for branded generic medicines based on current information.

"They have stated that they will continue to observe developments in the statutory and voluntary schemes during 2015 as more PPRS data becomes available. We are still concerned that scope for reduction remains an option and that the response does not adequately reflect the importance and operation of the branded generic medicines market in the UK. Branded generic medicines frequently offer additional patient benefits delivered through incremental innovation undertaken by the generic medicines industry.

"They already offer a big reduction off the price of the originator brand even after patent expiry and any further price reduction threatens the sustainability of this market and therefore the supply of these important medicines to patients."

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Notes for Editors:

The British Generic Manufacturers Association represents the interests of UK-based manufacturers and suppliers of generic medicines and promotes the development and understanding of the generic medicines industry in the United Kingdom.

Generic medicines contain the same active ingredient and are as effective as the equivalent brand and cost much less, making the NHS drugs bill affordable. More than two thirds of all medicines dispensed by the NHS are generics yet they cost only around 29% of the NHS drugs bill, a saving of more than £12.5billion in England & Wales alone. Without generics, the NHS drugs bill would be approximately twice its current level.

We represent the views and interests of our members and industry to the UK government, the devolved administrations, regulators, other relevant third parties, including where appropriate the Institutions of the European Union.