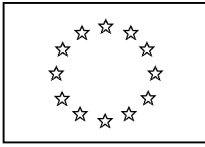


Brussels 18/06/2007 mj

**TO: MEMBERS OF THE STANDING COMMITTEE ON  
MEDICINAL PRODUCTS FOR HUMAN USE**

<b>Mr. R. SCHLOGEL (Austria)</b>
<b>Ms D. POULSEN (Denmark)</b>
<b>Ms E. METSÄRANTA (Finland)</b>
<b>Mr. J. MARIMBERT (France)</b>
<b>Prof. K. WOODS (United-Kingdom)</b>
<b>Dr. C. MORAITI (Greece)</b>
<b>Mr J.A. LISMAN (The Netherlands)</b>
<b>Pr. V. MARIA / Ms L. CARVALHO (Portugal)</b>
<b>Mrs. AM. A. NILSSON (Sweden)</b>
<b>Dr. A. LHOIR (Belgium)</b>
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<b>Mr. T. McGUINN (Ireland)</b>
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<b>Ms Z. ZEIFERTE (Latvia)</b>
<b>Mr R. MAČIULAITIS (Lithuania)</b>
<b>Mr A. KLIŠ / Ms M. SZELACHOWKA / Ms. A. OKULICH-KOZARYN (Poland)</b>
<b>Mr S. PRIMOŽIČ (Slovenia)</b>
<b>Dr. P. VELLA BONANNO /Ms. H. VELLA /Dr. J.J. BORG (Malta)</b>
<b>Ms D. STARÁ (Slovak Republic)</b>
<b>Dr R. ANCUCEANU, Dr R. BADESCU (Romania)</b>
<b>Mrs I. ATANASOVA, / Mrs S. SAPSSOVA (Bulgaria)</b>
<b>EMEA – Dr N. WATHION</b> <b>Vanessa Seguin (vanessa.seguin@emea.europa.eu)</b>
<b>Exelgyn Bipharma B.V.</b> <b>Regulatory Affairs, Catherine Denicourt (catherine.denicourt@exelgyn.com)</b>
<b>From:</b>
<b>Mr. Martin TERBERGER (EC)</b>

Cover plus one page



EUROPEAN COMMISSION  
DIRECTORATE GENERAL  
ENTERPRISE

**Commission Decision concerning the marketing authorization for the medicinal product for  
human use**

**"Mifegyne - mifepristone" (EMA/H/A-31/693)**

- The Commission adopted the decision concerning "**Mifegyne - mifepristone**", **(EMA/H/A-31/693), Art. 31 on 14<sup>th</sup> June 2007.**
- It will be notified forthwith to the Marketing Authorization holder.
- The attention of the national competent authorities has to be drawn to the fact that under EC Treaty general rules, a decision is a legal act whose publication is not obligatory in order to be binding.

Martin Terberger  
Head of Unit

*Pp Monika JAROS-LANQUETOT*