



Dr Sotirios G. Stergiopoulos appointed as Ipsen Chief Medical Officer

Paris (France), 26 January 2018 – Ipsen (Euronext: IPN; ADR: IPSEY) has announced that Sotirios G. Stergiopoulos, MD, has been appointed as Chief Medical Officer. Dr Stergiopoulos joined Ipsen in January 2017 as Senior Vice President, Head of Global Medical Affairs (GMA) and will retain this position in addition to the role as the new Chief Medical Officer within the company. Dr Stergiopoulos reports to Alexandre Lebeaut, MD, Executive Vice-President R&D and Chief Scientific Officer.

Dr Lebeaut commented, "Sotirios has extensive experience in directing global medical affairs strategies and a strong expertise in oncology drug development that includes chemotherapy, immunology drugs and targeted agents across various tumor indications. We are delighted to appoint him to an expanded role through which he will make the voice of the patient heard at the highest levels of the organization and represent the company externally as its primary medical representative."

Prior to joining Ipsen, Sotirios was Vice President, Head of Global Medical Affairs Oncology at Baxalta (now Shire, Cambridge, MA), Executive Medical Director Oncology US Medical Affairs at Celgene Corporation (Summit, NJ), Senior Global Brand Medical Director Oncology at Novartis Pharmaceuticals (East Hanover, NJ) and Director Medical Affairs Oncology at Bayer Healthcare (Montville, NJ).

Dr. Stergiopoulos added, "I am delighted to take on this leading role at Ipsen. This is an exciting time for Ipsen with significant growth and evolution. Together with our experienced leadership team, I look forward to helping our company continue to bring innovative new medicines to our patients."

Dr. Stergiopoulos is a physician executive with significant experience in the Pharmaceutical/Biotech industry, especially in Oncology. He has held appointments as an Attending Physician and trainee in institutions such as Albert Einstein College of Medicine, Harvard Medical School and the National Institutes of Health. He holds a Masters in Biotechnology Enterprise and Entrepreneurship (MBEE) from The Johns Hopkins University and a Medical Degree from Poznan University of Medical Sciences (Poland). Sotirios is a Fellow of the American College of Physicians, the New York Academy of Medicine as well as the Royal Society of Medicine (UK). He is also a Member of the American Association for Cancer Research and of the American Society of Clinical Oncology.

In October 2017 Dr. Stergiopoulos was appointed President of the Board of Governors for the Accreditation Council for Medical Affairs (ACMA); a body whose mission is to establish, certify, and maintain the competencies of qualified medical and scientific professionals who have a focus in Medical Affairs within the pharmaceutical & biotechnology industries.



About Ipsen

Ipsen is a global specialty-driven biopharmaceutical group focused on innovation and specialty care. The group develops and commercializes innovative medicines in three key therapeutic areas - Oncology, Neurosciences and Rare Diseases. Its commitment to oncology is exemplified through its growing portfolio of key therapies for prostate cancer, neuroendocrine tumors, renal cell carcinoma and pancreatic cancer. Ipsen also has a well-established Consumer Healthcare business. With total sales close to €1.6 billion in 2016, Ipsen sells more than 20 drugs in over 115 countries, with a direct commercial presence in more than 30 countries. Ipsen's R&D is focused on its innovative and differentiated technological platforms located in the heart of the leading biotechnological and life sciences hubs (Paris-Saclay, France; Oxford, UK; Cambridge, US). The Group has about 5,100 employees worldwide. Ipsen is listed in Paris (Euronext: IPN) and in the United States through a Sponsored Level I American Depositary Receipt program (ADR: IPSEY). For more information on Ipsen, visit www.ipsen.com.

Ipsen Forward Looking Statement

The forward-looking statements, objectives and targets contained herein are based on the Group's management strategy, current views and assumptions. Such statements involve known and unknown risks and uncertainties that may cause actual results, performance or events to differ materially from those anticipated herein. All of the above risks could affect the Group's future ability to achieve its financial targets, which were set assuming reasonable macroeconomic conditions based on the information available today. Use of the words "believes," "anticipates" and "expects" and similar expressions are intended to identify forward-looking statements, including the Group's expectations regarding future events, including regulatory filings and determinations. Moreover, the targets described in this document were prepared without taking into account external growth assumptions and potential future acquisitions, which may alter these parameters. These objectives are based on data and assumptions regarded as reasonable by the Group. These targets depend on conditions or facts likely to happen in the future, and not exclusively on historical data. Actual results may depart significantly from these targets given the occurrence of certain risks and uncertainties, notably the fact that a promising product in early development phase or clinical trial may end up never being launched on the market or reaching its commercial targets, notably for regulatory or competition reasons. The Group must face or might face competition from generic products that might translate into a loss of market share. Furthermore, the Research and Development process involves several stages each of which involves the substantial risk that the Group may fail to achieve its objectives and be forced to abandon its efforts with regards to a product in which it has invested significant sums. Therefore, the Group cannot be certain that favourable results obtained during pre-clinical trials will be confirmed subsequently during clinical trials, or that the results of clinical trials will be sufficient to demonstrate the safe and effective nature of the product concerned. There can be no guarantees a product will receive the necessary regulatory approvals or that the product will prove to be commercially successful. If underlying assumptions prove inaccurate or risks or uncertainties materialize, actual results may differ materially from those set forth in the forward-looking statements. Other risks and uncertainties include but are not limited to, general industry conditions and competition; general economic factors, including interest rate and currency exchange rate fluctuations; the impact of pharmaceutical industry regulation and health care legislation; global trends toward health care cost containment; technological advances, new products and patents attained by competitors; challenges inherent in new product development, including obtaining regulatory approval; the Group's ability to accurately predict future market conditions; manufacturing difficulties or delays; financial instability of international economies and sovereign risk; dependence on the effectiveness of the Group's patents and other protections for innovative products; and the exposure to litigation, including patent litigation, and/or regulatory actions. The Group also depends on third parties to develop and market some of its products which could potentially generate substantial royalties; these partners could behave in such ways which could cause damage to the Group's activities and financial results. The Group cannot be certain that its partners will fulfil their obligations. It might be unable to obtain any benefit from those agreements. A default by any of the Group's partners could generate lower revenues than expected. Such situations could have a negative impact on the Group's business, financial position or performance. The Group expressly disclaims any obligation or undertaking to update or revise any forward looking statements, targets or estimates contained in this press release to reflect any change in events, conditions, assumptions or circumstances on which any such statements are based,



unless so required by applicable law. The Group's business is subject to the risk factors outlined in its registration documents filed with the French Autorité des Marchés Financiers.

The risks and uncertainties set out are not exhaustive and the reader is advised to refer to the Group's 2016 Registration Document available on its website (www.ipsen.com).

For further information:

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