

**Ipsen appoints Steven Hildemann, MD., PhD
as Executive Vice President, Chief Medical Officer**

PARIS, France, 14 January 2020 -- Ipsen (Euronext: IPN; ADR: IPSEY) today announced the appointment of Dr. Steven Hildemann as Executive Vice President, Chief Medical Officer, Head of Global Medical Affairs and Pharmacovigilance effective March 1, 2020. Based in Paris, France, Dr. Hildemann will report directly to Aymeric Le Chatelier, CEO, Ipsen and serve on the Executive Leadership Team.

Aymeric Le Chatelier, CEO, Ipsen, commented, *“As we advance as a leading global biopharmaceutical company focused on innovation and Specialty Care, it is my great pleasure to appoint Dr. Hildemann to lead Ipsen’s Global Medical Affairs, Patient Affairs and Pharmacovigilance organizations through patient-centric leadership, sound medical governance and business conduct. With over 20 years of service in the pharmaceutical industry and 10 years as a physician-scientist in academic medicine, he brings a wealth of experience in medical affairs, clinical operations and patient safety from major global biopharmaceutical companies. Dr. Hildemann will play a crucial role in delivering on our global medical and patient safety strategy, engaging in structured, ethical and high-quality dialogue with patients, external and internal stakeholders throughout the entire life-cycle of Ipsen’s portfolio.”*

Dr. Hildemann will act as a strategic partner to Ipsen’s scientific, clinical and business teams. He will provide a global medical strategy based on real value to patients and caregivers. In close partnership with R&D, Dr. Hildemann will lead the generation of post-launch clinical trial data by defining, prioritizing, optimizing or guiding studies in line with this strategy. As a member of the Executive Leadership team, he will actively contribute to the overall management and strategic leadership of the company.

Prior to joining Ipsen, Dr. Hildemann held leadership roles in science-based bioethics and built an innovative digital health startup in cancer care after serving for five years as Chief Medical Officer, Senior Vice President, Head of Global Medical Affairs and Global Patient Safety at Merck. He also held several strategic leadership positions with biopharmaceutical companies such as Pharmacia-Pfizer and Schering-Plough-MSD. Dr. Hildemann is board certified in internal medicine and cardiology with broad clinical training across internal medicine including medical oncology, gastroenterology, rheumatology and pulmonary oncology at university hospitals in Munich, Germany. Throughout his career, he has engaged in part-time clinical practice, late-stage pharmaceutical research and medical teaching. Dr. Hildemann received his MD-PhD at the Albert Ludwig University of Freiburg, Germany, where he continues to serve as an adjunct Professor of Medicine.

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, Neuroscience 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 over €2.2 billion in 2018, 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 5,800 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.fr

Ipsen—Cautionary Note Regarding Forward-Looking Statements

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 favorable 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 2018 Registration Document available on its website (www.ipsen.com).

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