

## **Ad hoc announcement**

### **Inside information pursuant to Article 17 MAR**

#### **Heidelberg Pharma AG Announces Adjustment of Guidance**

**Ladenburg, Germany, 4 October 2022** – Heidelberg Pharma AG (FSE: HPHA) today announced that it has adjusted its guidance for the current fiscal year published on 24 March 2022. The license agreement with the strategic partner Huadong Medicine Co., Ltd., Hangzhou, China, and the corresponding license payment increased Heidelberg Pharma's sales revenue significantly. Development expenses remained below planning due to the later production of intermediate steps for the follow-on candidates. Both factors have an influence on the operating result that will improve and funding requirements for fiscal year 2022 will decrease.

The Heidelberg Pharma Group expects for the financial year 2022 sales and other income between EUR 18.5 million and EUR 20.5 million (previously: EUR 7.5 million to EUR 9.5 million). Operating expenses will range between EUR 35.0 million and EUR 39.0 million (previously: EUR 41.0 million to EUR 45.0 million). Based on these adjustments, an operating result (EBIT) between EUR -16.0 million and EUR -20.0 million is expected (previously: EUR -32.5 million to EUR -36.5 million).

For 2022, Heidelberg Pharma anticipates cash requirements of EUR 8.0 million to EUR 11.0 million (previously: EUR 33.0 million to EUR 37.0 million). Monthly cash consumption is expected to range between EUR 0.6 million and EUR 0.9 million per month (previously: EUR 2.8 million and EUR 3.1 million). Based on the existing planning and the capital increase that was closed after the end of the reporting period, the company assumes a financing range until mid-2025.

The Interim Management Statement on the first nine months of 2022 will be published as planned on 13 October 2022.

+++ End of the ad hoc announcement +++

#### **About Heidelberg Pharma**

Heidelberg Pharma is an oncology specialist and the first company to develop the toxin Amanitin into cancer therapies using its proprietary ATAC<sup>®</sup> technology and to advance the biological mode of action of the toxin as a novel therapeutic principle. The proprietary technology platform is being applied to develop the Company's own therapeutic ATACs<sup>®</sup> as well as in third-party collaborations. The lead candidate HDP-101 is a BCMA ATAC in clinical development for multiple myeloma. HDP-102, a CD37 ATAC for Non-Hodgkin lymphoma and HDP-103, a PSMA ATAC for metastatic castration-resistant prostate cancer, are in preclinical testing.

Heidelberg Pharma AG is based in Ladenburg, Germany, and is listed on the Frankfurt Stock Exchange: ISIN DE000A11QVV0 / WKN A11QVV / Symbol HPHA. More information is available at <http://www.heidelberg-pharma.com/>.

ATAC<sup>®</sup> is a registered EU trademark of Heidelberg Pharma Research GmbH.

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