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CSPC PHARMACEUTICAL GROUP LIMITED

石藥集團有限公司

(Incorporated in Hong Kong with limited liability)

(Stock Code: 1093)

VOLUNTARY ANNOUNCEMENT

DATA FROM PHASE I CLINICAL STUDY OF SYS6002 AS MONOTHERAPY FOR THE TREATMENT OF ADVANCED CERVICAL CANCER PRESENTED AS A RAPID ORAL PRESENTATION AT THE 2026 ANNUAL GLOBAL MEETING OF THE INTERNATIONAL GYNECOLOGIC CANCER SOCIETY (IGCS 2026)

The Board of Directors (the “**Board**”) of CSPC Pharmaceutical Group Limited (the “**Company**”, together with its subsidiaries, the “**Group**”) is pleased to announce that the Group has presented the data from the Phase I clinical study (the “**Study**”) of SYS6002 as monotherapy for the treatment of advanced cervical cancer as a rapid oral presentation at the 2026 Annual Global Meeting of the International Gynecologic Cancer Society (IGCS 2026).

OVERVIEW OF THE STUDY

The Study is a nationwide, multi-centre Phase I clinical trial designed to evaluate the safety, tolerability, pharmacokinetic profile and preliminary efficacy of SYS6002 in patients with advanced solid tumours. At this meeting, the Group presented the trial results of SYS6002 as monotherapy for the treatment of previously treated advanced cervical cancer.

As at 8 May 2026, a total of 82 patients were enrolled in the Study and assigned to two dose cohorts, including 39 patients in the 2.4 mg/kg once every 2 weeks (“**Q2W**”) cohort and 43 patients in the 3.6 mg/kg once every 3 weeks (“**Q3W**”) cohort, with a median follow-up of 16.1 months. The relevant safety and efficacy data are as follows:

- In terms of safety, the safety profile of the above two dose cohorts was generally manageable, with treatment-related adverse events mostly being Grade 1 to 2. Among them, ocular-related adverse events were the most common, but no Grade 3 or above events occurred, and no patient discontinued treatment due to these events. The incidence of interstitial lung disease and non-infectious pneumonitis was low (2.4%). The incidence and severity of treatment-related rash were both low (any grade: 20.7%; Grade 3 or above: 1.2%), and no cases of Stevens-Johnson syndrome

or toxic epidermal necrolysis were reported. The incidence and grade of peripheral neuropathy were also low (12.2%; all Grade 1 to 2). In addition, no treatment-related adverse events leading to death occurred.

- In terms of efficacy, both dosing regimens demonstrated preliminary efficacy in the study population. The confirmed objective response rate (ORR) was 35.9% in the 2.4 mg/kg Q2W cohort and 28.6% in the 3.6 mg/kg Q3W cohort. The median duration of response (DoR) for patients who achieved a confirmed response in the two cohorts was 8.2 months and 10.3 months, respectively; the median progression-free survival (PFS) was 4.1 months and 4.3 months, respectively; and the median overall survival (OS) was 15.1 months and 11.5 months, respectively.

Currently, the study of SYS6002 2.4 mg/kg Q2W regimen for the treatment of advanced cervical cancer previously treated with immunotherapy has entered a Phase III clinical trial (NCT07230626).

ABOUT SYS6002

SYS6002 is a next-generation antibody-drug conjugate (ADC) targeting Nectin-4. By utilising enzyme-catalysed site-specific antibody conjugation technology, the drug specifically directs the potent mitotic inhibitor MMAE to Nectin-4-expressing cancer cells. In addition, the stable linker design of SYS6002 not only facilitates the delivery of high concentrations of MMAE to the tumour but also reduces toxicity by decreasing systemic exposure to the toxin.

By order of the Board
CSPC Pharmaceutical Group Limited
CAI Dong Chen
Chairman

Hong Kong, 8 October 2026

As at the date of this announcement, the Board comprises Mr. CAI Dong Chen, Dr. CAI Lei, Mr. WEI Qingjie, Mr. ZHANG Cuilong, Mr. WANG Zhenguo, Dr. LI Chunlei, Dr. YAO Bing, Mr. CAI Xin, Mr. CHEN Weiping, Mr. QU Zhiyong and Mr. ZHANG Yiwei as Executive Directors; and Mr. WANG Bo, Mr. CHEN Chuan, Prof. WANG Hongguang, Mr. AU Chun Kwok Alan, Mr. LAW Cheuk Kin Stephen and Ms. LI Quan as Independent Non-executive Directors.