**Appendix No. 2 to Request for Bids No. 01/WPD101/2020**

#### ..................................., on ...........................

*(place) (date)*

**CONTRACTOR**

……………………………………….

*(name/registered office/address/tax ID (NIP) of the Contractor)*

**AWARDING ENTITY:**

WPD Pharmaceuticals sp. z o. o.

ul. Żwirki i Wigury 101,

02-089 Warszawa,

#### **DECLARATION OF COMPLIENCE WITH THE REQUIREMENTS INDICATED IN SECTION VI OF THE REQUEST FOR BIDS**

For the purpose of the contract award procedure covering research service involving process and analytical development, non-GMP and GMP manufacture of drug substance, for consistent generation of the biopharmaceutical recombinant-protein product, as a part of the project No. POIR.01.01.01-00-0912/17, entitled: “*Development of a new drug used in the therapy of glioblastoma multiforme”* (Request for Bids No. 01/WPD101/2020), on behalf of the Contractor I hereby declare that

Company …………………………………………………………… *(Company name)*

Registered in …………………………………………………………*(Registered Address)*

meets the following conditions:

1. I declare that we have adequate knowledge and experience within the contract object and people declare that all data contained in the Bid capable of performing the order.
2. I declare that we are in financial and economic situation that will allow the contract to be performed with due diligence.
3. I declare that:
4. we have at least (2) two orders duly completed, in particular consisting of generation and maintenance and analysis of microbial cell banks and manufacturing of recombinant protein products according in compliance with GMP standard;
5. we have and engage to perform the contract persons with appropriate qualification and experience, including at least (2) persons with a specialized education (in pharmacy, chemistry, biotechnology or related fields, i.e. medicine, biological sciences), experienced in recombinant protein production with the use of microbial and insect cell expression systems in GMP compliant standard and process optimization and characterization of the expression systems and analytical method development and validation;
6. we have appropriate technical facility and optimal equipment dedicated to the production, maintenance and testing of the microbial cell expressions systems/banks, as well as manufacturing, maintenance and testing of the recombinant protein products obtained with the use of the microbial expression systems in compliance with GMP, and have certificate of the GMP standards for the production of the above-mentioned as API or have other required by law documents for the production, testing and storage of the cell banks and the protein based API, such as valid authorization for API manufacturing for the purpose of clinical trials.
7. I declare that all data contained in the Bid are consistent with the actual and legal status.

**Curriculum vitae of the persons participating in the contract object preparation is attached to the Bid.**

Place ………………, on ……….… ……..................................................

*(signature of the person(s) authorized to represent the Contractor)*