



May 2024

General Terms and Conditions for Genomic Services

1. GENERAL

- a. These terms and conditions ("Agreement") apply to all genomic services ("Services") performed by QIAGEN Genomic Services ("QIAGEN") to its customers (each, a "Customer"), unless QIAGEN and Customer have agreed to a separate written contract for the same purpose and scope signed by authorized representatives of both parties (a "Contract"). To the extent the parties have executed a Contract, and there are any conflicts between these terms and those in the Contract, then the Contract shall control. Any provisions printed or otherwise contained in any purchase order, acknowledgement, acceptance or other document from Customer purporting to govern the purchase of Services from QIAGEN which are inconsistent with or in addition to the Agreement shall have no force or effect and shall not constitute any part of the Agreement between the parties.
- b. The Services to be performed by QIAGEN shall be described in an agreed Sample Submission Form ("SSF"). The SSF shall include any and all Quote(s) related to said SSF. Upon formal SSF approval by QIAGEN, Customer prints, signs and provides the SSF to QIAGEN to initiate the Services. In case Samples are provided by Customer to QIAGEN, Customer adds the SSF to the Sample shipment. Customer adds the individual project number provided by QIAGEN to shipment address.
- c. Customer hereby acknowledges and agrees that QIAGEN performs Services both in QIAGEN's own laboratories ("QIAGEN Genomic Services Sites") and in cooperation with QIAGEN partners ("QIAGEN Partners"). QIAGEN assigns the Services to the respective service laboratories based on data integrity, capacity and expertise to provide high quality Services to the Customers. QIAGEN collaborates only with

contract research partners that comply with QIAGEN's corporate quality and supply chain standards.

- d. The Services are intended for molecular biology applications.

The Services are not intended for the diagnosis, prevention, or treatment of a disease.

2. DEFINITIONS

- a. "Licensed Materials" shall mean any and all (proprietary) software, databases, algorithms, materials, Processes and applications owned or licensed by QIAGEN that are used in the Services. Nothing in this Agreement shall be construed as granting to Customer, by implication, estoppel or otherwise, any license or other proprietary right with respect to any Licensed Material or to any intellectual property owned or controlled by or licensed to QIAGEN. For the avoidance of doubt, Licensed Materials does not encompass Results.
- b. "Results" shall mean any information, designs, technical drawings, algorithms, elaborated design data, experimental data, nucleic acid sequences, amino acid sequences, compositions created and tested in the lab, technical or industrial data, tools, processes, methodology, and any intellectual property therein, created by QIAGEN in the course of the Services and provided to the Customer under the present Agreement.
- c. "Customer Biological Data" shall mean any and all data, documents, know-how, analyses and information, whether in writing, orally or in any other form, provided or disclosed by Customer or a third party on behalf of Customer to QIAGEN in relation to the Services performed under the present Agreement.
- d. "Samples" shall mean any and all physical material provided by Customer under the present Agreement.
- e. "Process(es)" shall mean any operation/workflow which is performed utilizing Samples and/or Customer Biological Data to complete the Project defined in the SSF, whether or not by automatic (including, without limitation, accessing, collecting, recording, organizing, retaining, storing, adapting or altering, transmitting,

retrieving, consulting, using, aligning, combining, extrapolating, analyzing, interpreting information) or any other means.

- f. "Support" shall mean any and all assistance, consultation and guidance provided by QIAGEN to Customer with regard to the Project as defined in the SSF.
- g. "Project shall mean the scope of work on Samples and Customer Biological Data as agreed upon between QIAGEN and Customer in the SSF.

3. PRICES AND PAYMENT TERMS

- a. QIAGEN shall be entitled to increase the prices at any time with thirty (30) calendar days' notice.
- b. Unless otherwise specified in writing by QIAGEN or by virtue of law, the prices are exclusive of transportation, insurance, license fees, customs duties, withholding, value added tax and any sales, use, excise, and other similar taxes. Customer shall pay all such fees, duties, and taxes in addition and in the manner and at the rate prescribed by the relevant authority or reimburse QIAGEN for all federal, state or local sales, use or other taxes, fees or duties arising out of their agreement or the transactions contemplated by their agreement, if any (other than taxes based on QIAGENs net income).
- c. The full amount of each invoice for Services hereunder shall be paid net 30 days (terms contingent upon credit worthiness of Customer) from the date of the invoice unless otherwise stated on the face of the invoice.
- d. QIAGEN reserves the right to assess a late fee equal to one and one-half percent (1.5%) per month or, if lower, the maximum amount permitted by applicable law, on all amounts not paid when due, calculated on a daily basis beginning with the 1st day following the invoice due date. Any check or remittance received from or for the account of Customer may be accepted and applied by QIAGEN against any indebtedness owing by Customer, without prejudice to, or the discharge of, the remainder of any such indebtedness regardless of any condition, provision, statement, legend or notation appearing on, referring to or accompanying any check or remittance.

4. SAMPLE AND MATERIAL DELIVERY

- a. Customer shall supply QIAGEN with Samples in the quantities and quality and on time as specified in the SSF.
- b. Customer is responsible for the delivery of Samples to QIAGEN in such a manner as to maintain integrity, and in accordance with any relevant shipping and handling regulations or instructions and related costs. Customer is required to get in contact with QIAGEN before shipping the Samples to clarify potential customs issues (documents, permits, etc.). QIAGEN shall be entitled to carry out an inspection upon delivery of Samples in order to determine their condition prior to processing the Samples. If the result of said inspection confirms that processing of Samples is impossible or impossible under standard conditions (by way of example due to but not limited to impurities, contaminations, deterioration, decomposition, or the like) QIAGEN will not process the Samples and inform Customer accordingly. If processing of the Samples is only possible with increased effort (e.g. non-standard protocols), QIAGEN will provide Customer with a respective quote for additional costs. Upon payment of costs by Customer QIAGEN will continue processing the Samples.
- c. Customer will assure, to the best of their knowledge, that the Samples are free of any substance, virus, bacteria etc. harmful to personnel. Any pathogen or living organism (e.g. bacteria) which is not considered to be inactivated needs to be announced before shipping and Customer must wait for approval before shipping. This is also true for any living non-human organism and/or non-human sample material sent from outside of the EU. If applicable, Customer will provide QIAGEN with relevant occupational safety information in the possession of Customer. Customer shall also announce all Samples from endangered species prior to shipment and shall provide appropriate documents and regulatory approvals for the processing and shipment of endangered species Samples to QIAGEN.
- d. Customer shall bear the costs for shipping the Samples to QIAGEN Genomic Services Sites. Shipment will be effected by Customer's preferred logistics company.
- e. Customer will not provide any person-identifiable data to identify Samples. Customer shall not transmit or disclose in any way to QIAGEN any personal health information

as defined by applicable regulations. By ordering, Customer confirms to have acquired all the required ethical permissions and regulatory approval for the proposed work and QIAGEN will have no liability if these permissions and approvals have not been properly granted.

- f. QIAGEN shall make no representations or warranties with respect to the Samples in connection with the performance of the Services. Customer agrees to indemnify QIAGEN from all actions and liabilities resulting from QIAGEN's processing of any Samples, Customer Biological Data or the use of any Results produced by QIAGEN in the course of the Project.
- g. QIAGEN acknowledges that Customer Biological Data and Samples are and shall remain the exclusive property of Customer, and that nothing in this Agreement shall be construed as granting to QIAGEN, by implication, estoppel or otherwise, any license or proprietary right with respect to such Customer Biological Data or Samples unless necessary to carry out the Services.
- h. QIAGEN agrees not to use Customer Biological Data for any other purpose than the performance of the Services. QIAGEN agrees not to analyze, chemically modify, use, reverse engineer or determine the structure of any Samples unless necessary to carry out the Services.

5. SAMPLE STORAGE

Any remaining Samples shall either be returned to Customer upon request or discarded after three months from finalization of Services according to the SSF. Customer is responsible for the import of these Samples from QIAGEN Genomic Services Sites and bears any import related costs incurred.

6. DATA, RESULTS, AND SUPPORT

- a. The Results generated during or resulting directly from the Services performed by QIAGEN shall be the exclusive property of Customer. Results will be discarded after 90 days from finalization of Services according to the SSF. QIAGEN will keep all information confidential and will not exploit Results for any use other than general statistic purposes, troubleshooting and improvements of Services.
- b. Delivery of Results. Customer agrees that Results will be shared via QIAGEN's internal servers, cloud-based servers or via hard drive delivery, all at QIAGEN's sole discretion.
- c. Customer Biological Data and security. As between the parties, Customer and its supplier(s) own and shall retain title to all intellectual property rights and other proprietary rights in and to the Customer Biological Data provided by Customer or Customer representatives. Customer grants QIAGEN the right to reproduce, adapt, distribute, publish, use, and share the Customer Biological Data solely for the purposes of (i) processing Customer's requests, transactions, and analyses contemplated herein, and (ii) for internal purposes. QIAGEN agrees to maintain commercially reasonable security procedures with respect to access and storage and sharing of the Customer Biological Data. These procedures are intended to provide reasonably appropriate technical and organizational safeguards against unauthorized disclosure or access. QIAGEN has no obligation to maintain access to Customer Biological Data and may delete Customer Biological Data from its systems at any time.
- d. Licensed Materials. Customer acknowledges that QIAGEN and its supplier(s) own and shall retain all intellectual property rights and other proprietary rights in and to the Licensed Materials and any other materials and information QIAGEN provides to Customer as part of this Agreement, including without limitation any derivatives, improvements or modifications of the foregoing, whether or not made by QIAGEN. Customer Biological Data are not considered Licensed Materials. For clarity, any discoveries that Customer makes while using the Results are not considered to be QIAGEN intellectual property.

- e. Adverse Actions. Customer hereby acknowledges QIAGEN's ownership and rights in the Licensed Materials and Processes. To the extent legally enforceable in the jurisdiction relevant to the Licensed Materials and/or Processes in issue, Customer and its affiliates shall not participate as an adverse party in, or otherwise provide material support to, any legal action, litigation, arbitration, mediation, opposition, re-examination, revocation, nullity proceeding or other legal or administrative proceeding anywhere in the world that (i) challenges the enforceability, scope, validity, or essentiality or seeks to determine the value or construction of any patent of the Licensed Materials and/or Processes or part thereof, or (ii) alleges unfair competition or patent misuse involving the Licensed Materials and/or Processes. In the event Customer or any of its affiliates actively participates as an adverse party in, or otherwise provides material support to, any such action, unless all claims of all Licensed Materials and/or Processes involved in the action have been declared invalid, Customer shall pay all of QIAGEN's costs associated with the action, including without limitation travel and attorney's fees.
- f. Feedback. To the extent Customer (and/or Customer representatives) provide or make available to QIAGEN any suggestions; ideas; improvements; modifications; feedback; error identifications; content corrections or additions; variant classifications; pooled anonymized allele frequency data, opinions regarding the appropriateness of a particular curated article to the clinical assessment of one or more variants; or other content or information related to the Licensed Materials (other than the Customer Biological Data) ("Feedback"), Customer and Customer representatives hereby grant QIAGEN a fully paid-up, irrevocable, perpetual, worldwide, nonexclusive license, with full rights to sublicense, to: (i) use and exploit such Feedback to improve QIAGEN's Services and, (ii) use, reproduce, prepare derivative works of, perform, display, make, sell and otherwise distribute Services incorporating or utilizing such Feedback.
- g. De-identified Data Use. QIAGEN may utilize data capture, syndication and analysis tools, and other similar tools, to create, extract, compile, keep, aggregate or synthesize data, usage patterns or information which has been de-identified consistent with applicable data privacy laws and associated data protection standards

contained in Customer Biological Data (“De-identified Data”). QIAGEN may (i) use and analyze the De-identified Data internally to test, develop and improve QIAGEN’s Services, (ii) use usage patterns as part of QIAGEN’s Services, and (iii) use De-identified Data obtained through QIAGEN data analysis pipelines and knowledgebases for commercial use and other applications, including without limitation, health, research or patient care. QIAGEN will ensure that no personally identifiable information provided by Customer is disclosed in the course of the QIAGEN Services to any third party without Customer’s consent.

- h. Support. Consultation and support will be provided for 90 days following delivery of data (for data delivery only projects), or delivery of data analysis (for data analysis inclusive projects). For extended support beyond 90 days, additional costs will be at the Customer’s expense.
- i. QIAGEN gives no representation or warranty that the Results and/or the use(s) thereof do not infringe the intellectual property or any other rights of third parties unless expressly stated in the present Agreement.
- j. To the extent the Samples are Personal Data (as defined in the applicable data protection laws) the parties will comply with all applicable data protection laws including without limitation Regulation (EU) 2016/679 in relation the processing of the Personal Data disclosed under this Agreement and shall execute the data processing agreement. QIAGEN will not (i) process or otherwise use Personal Data other than as necessary to perform this Agreement or as otherwise directed by Customer or (ii) disclose such Personal Data to any third party, unless otherwise permitted hereunder.

If such Personal Data from the European Economic Area (EEA), the United Kingdom and Switzerland is processed by QIAGEN entity based outside of these territories, the EU-US and/or Swiss-US Privacy Shield, and/or the Standard Contractual Clauses shall apply. To the extent that the California Consumer Privacy Act (“CCPA”) is applicable to either party such party agrees to comply with all of its obligations under the CCPA, including but not limited to: (i) not to sell the Personal Data; (ii) retain, use, or disclose the Personal Data for any purpose other than for the specific purpose of performing the services; (iii) retain, use, or disclose the Personal Data for a commercial purpose

other than providing the services; or (iv) retain, use, or disclose the information outside of the direct business relationship between the parties.

7. DELIVERY DATES, COMPLETION PERIOD

QIAGEN shall endeavor to fulfil Services as agreed upon in the SSF in good time. The processing time is defined as number of working days from receipt of Customer Biological Data and/or Samples and SSF at QIAGEN to the day on which Results are provided to the Customer. Closed days, public holidays, Saturdays and Sundays are not considered working days. QIAGEN is entitled to make partial deliveries. Delays due to force majeure, interruption of operations or other circumstances for which QIAGEN is not responsible shall release them from their obligation to perform for the duration of the disruption or the effects of the disruption. Claims for damages by Customer are excluded in this case. If Results are provided late, Customer shall be obliged to grant QIAGEN a reasonable grace period. Only after the expiry of the grace period of 3 weeks may Customer resign from the SSF if QIAGEN has not yet provided any Results for reasons for which QIAGEN is not responsible. Where Customer decides to resign from the SSF, QIAGEN will invoice Customer for the Services performed to date and, where applicable, refund to Customer the difference to any payments already made. In such case all further clauses of the Agreement will remain in force.

8. TRANSFER OF TITLE AND OTHER RIGHTS

- a. Any and all discoveries, inventions, ideas, know-how, developments, formulas, techniques, data and any other information derived or generated by Customer from the Results shall be the exclusive property of Customer. However, nothing in this Agreement shall constitute a grant of license to Customer to commercially exploit any of QIAGEN's technologies, intellectual property or other rights for any purpose whatsoever.
- b. Customer acknowledges that QIAGEN possesses certain expertise, know-how, techniques, processes, intellectual property and other rights, which have been independently developed by QIAGEN, and which relate to QIAGEN's business

operations in general (hereinafter the "QIAGEN Property"). QIAGEN Property is and remains the exclusive property of QIAGEN and QIAGEN shall not be restricted in using QIAGEN Property. Nothing in this Agreement shall be construed as granting to Customer, by implication, estoppel or otherwise, any license or other access to QIAGEN Property.

- c. QIAGEN shall not be entitled to publish any Results, unless otherwise agreed upon in the SSF.

9. LIMITED WARRANTIES

- a. The parties agree that Services shall be deemed accepted if Customer does not give notice to the contrary within a period of one week from receipt of Results. In case Results are not accepted by Customer, Customer will contact QIAGEN immediately and provide a statement of grounds for not accepting the Results. In case Results are accepted, Customer is solely responsible for any legal or other consequences resulting from relying on or use of Results.
- b. If Customer contests the correctness of the Results according to Sec. 9.1 of the present Agreement, Customer shall not be entitled to withhold payment unless the defectiveness of the Results are (i) acknowledged by QIAGEN or (ii) subject to a final court decision.
- c. Customer is responsible for the proper delivery of the Samples. QIAGEN is not liable for the loss or damage of any Samples during transport. Customer shall be exclusively and at all times liable for the safety, packaging and insurance of the Samples from the time of its dispatch until delivery to QIAGEN.
- d. Customer guarantees and undertakes to QIAGEN to ensure that all Samples sent to QIAGEN for the purpose of carrying out the Services are available in safe form. Customer further undertakes to indemnify QIAGEN and its personnel or other representatives against all losses, costs and other damages incurred by QIAGEN as a result of Samples being dangerous, unless this fact is not attributable to Customer. If Samples are dangerous, the contracting authority shall notify the supplier thereof in

writing before dispatch. Customer is also obliged to mark packaging, Samples and/or containers accordingly.

10. LIABILITY

- a. During the normal course of performing the Services, some experiments may fail to produce usable Results or Samples may accidentally be lost or damaged. QIAGEN assumes no liability for the loss of Samples provided that all usual and reasonable processes and safeguards have been observed. In case it is proven that loss or damage of Samples has occurred due to QIAGEN's non-observance of aforementioned processes and safeguards, QIAGEN shall reimburse Customer the costs of such Samples up to a maximum amount of the amounts received by QIAGEN for the Services within the last twelve (12) months. Results will vary depending on the quality of Samples and QIAGEN provides no warranty as to the quality of any Results generated, and shall not be held liable for any damage or cost incurred from the use of such Results.
- b. QIAGEN shall not be liable to Customer for any indirect, incidental, special or consequential loss or damages, including, without limitation, loss of profits, unless such liability is due to QIAGEN's negligence and/or willful misconduct. Save in respect of personal injury or death, QIAGEN's total liability for loss or damage arising out of or in relation to this Agreement shall in no event exceed the amount paid by Customer to QIAGEN with regard to this Agreement.
- c. QIAGEN shall not be obliged to pay compensation for delays, errors, damages or other problems caused by events or circumstances which were unforeseeable for QIAGEN or which are beyond its control or which result from compliance with official orders, laws or regulations.
- d. If QIAGEN is affected by substantial disadvantages (availability of materials to be used; change of the legal framework conditions) due to interruptions or delays in performance for which QIAGEN is not responsible, in particular scheduling difficulties, QIAGEN shall be entitled to resign from the SSF in whole or in part with regard to the part not yet performed. In the event QIAGEN resigns from this SSF,

QIAGEN will invoice Customer for the Services performed to date and refund to Customer the difference to any payments already made.

If the impediment lasts longer than 6 weeks, Customer shall be entitled to resign from the unfulfilled part of the SSF after the unsuccessful expiry of a reasonable grace period. Where Customer resigns from the unfulfilled part of the SSF in accordance with the above, QIAGEN will invoice Customer for the Services performed to date and, where applicable, refund to Customer the difference to any payments already made.

11. UNACCEPTED RESULTS

Complaints relating to Results may only be made in accordance with the rules set out in Sec. 9.1. In any case in which the inaccuracy of the Results is not amicably acknowledged by QIAGEN and Customer, Customer shall bear the cost for (i) repeating the Project or (ii) the necessary verification of the Results.

12. CONFIDENTIALITY

- a. Any and all Customer Biological Data and Samples, disclosed to or provided to QIAGEN under this Agreement including the SSF, shall be considered as confidential (hereinafter collectively referred to as "Confidential Information").
- b. QIAGEN shall use Confidential Information only for the performance of the Services under the present Agreement and shall not exploit, whether directly or indirectly, any Confidential Information for its own benefit or the benefit of any third party (person or entity) without the specific prior written consent of Customer. Any use of Confidential Information shall be in accordance with this Agreement and the SSF.
- c. QIAGEN shall maintain the Confidential Information in confidence and shall not disclose, directly or indirectly, Confidential Information to any third party (person or entity), other than its duly authorized representatives, employees, consultants and approved sub-QIAGENS who have a need to know such Confidential Information in the course of the performance of their duties relating to the Services. QIAGEN shall

advise all such persons who receive or are to receive Confidential Information that such information is confidential and may only be used for the Services, and shall require their compliance with the terms of this Agreement. QIAGEN shall maintain at least the same standard of custody of Confidential Information as QIAGEN keeps custody of the QIAGEN's own confidential information, but no less than a standard of care that is reasonable under the circumstances to maintain secrecy and control disclosure. QIAGEN shall immediately notify Customer if QIAGEN becomes aware of any suspected or actual unauthorized use, copying or disclosure of Confidential Information.

- d. The confidentiality and non-use of obligation of Sec. 12 of the present Agreement shall not apply to Confidential Information that:
- i. at the time of disclosure, is already in the public domain through no fault of QIAGEN;
 - ii. after disclosure, becomes part of the public domain by disclosure through no violation of this Agreement;
 - iii. QIAGEN is able to prove, has been lawfully in QIAGEN's possession prior to any disclosure under this Agreement;
 - iv. is hereafter lawfully disclosed by a third party to QIAGEN, where such third party did not acquire such information under a still effective obligation of confidentiality to Customer;
 - v. is required to be disclosed by an order or action of a governmental agency, authority or court (provided that Customer shall be informed as soon as reasonably possible and provided that QIAGEN shall furnish only that portion of the Confidential Information which is legally required, and shall exercise all efforts required to obtain confidential treatment for such information).
- e. If Confidential Information was disclosed by Customer to QIAGEN prior to the execution of this Agreement (as defined below) or prior to entering into an SSF under the present Agreement in anticipation of the parties entering into this Agreement or an SSF thereunder, such Confidential Information shall be treated as confidential and be subject to the terms and conditions of this Agreement.

- f. The obligations of confidentiality and non-use set forth herein shall remain in effect for a period of seven (7) years from the last date on which Confidential Information was disclosed to QIAGEN.
- g. Neither party may use the other party's name or trademark, or the name or trademark of any employee or agent, in any advertising or sales promotional material without the prior written consent of the other party.

13. INDEMNIFICATION

Customer shall indemnify, defend and hold harmless (collectively "Indemnify") QIAGEN, its affiliates and its and their respective directors, officers, employees, stockholders and agents (each, a "QIAGEN Indemnitee") from and against any and all losses, damages, liabilities, judgments, fines, amounts paid in settlement, expenses and costs of defense (including without limitation reasonable attorneys' fees and witness fees) ("Losses") resulting or arising from any claim, action, proceedings investigation or litigation brought or initiated by a third party ("Third Party Claim") to the extent that such Third Party Claim arises out of (a) the use, marketing, sale or distribution of the Results by Customer, (b) a violation of any law, convention or other regulation in any jurisdiction, including without limitation any intellectual property law or convention, by QIAGEN in the course of the performance of any and all Services under the present Agreement unless the violation is based on the Licensed Materials, or (c) the breach of this Agreement by Customer, except that Customer shall not Indemnify QIAGEN to the extent that any such Losses arise from the gross negligence or willful misconduct of any QIAGEN Indemnitee.

14. GENERAL COMPLIANCE

Each of the parties represents and warrants to the other party that it will comply with all applicable laws, rules or regulations (“Applicable Laws”), including, but not limited to, applicable federal, state and local laws, rules, regulations and guidelines relating to the manufacturing, quality control, packaging, labeling, handling, shipping, importation, exportation and storage of Samples.

15. TRADE COMPLIANCE

Customer shall comply with all applicable customs and export control regulations including but not limited to those issued by the United Nations, the US Government and the European Union. In particular, without limitation, Customer shall refrain from any transactions in relation to QIAGEN’s Services which would violate any applicable sanctions, embargoes or foreign trade restrictions including but not limited to those issued by the United Nations, the US Government and the European Union (referred to as “Sanctions”). Customer declares that it is not a person targeted by Sanctions nor is it otherwise owned or controlled by or acting on behalf of any person targeted by Sanctions. Customer shall promptly notify QIAGEN in writing if the Customer or any of the aforementioned persons become subject to Sanctions. Furthermore, the Customer commits to providing QIAGEN with all necessary documents and information required for Sanctions control checks performed by QIAGEN and/or any relevant authority. In the event of any violation of applicable Sanctions by the Customer, the Customer shall fully indemnify and hold harmless QIAGEN from and against any and all losses, damages, liabilities, costs, and expenses (including reasonable attorneys' fees) resulting from such violations.

16. NON-DISCRIMINATION

Neither Customer nor QIAGEN shall discriminate in the performance of this Agreement because of race, color, sex, sexual orientation, age, religion, handicap, marital status, or national origin in violation of any applicable federal, state, or local law or regulation.

17. BINDING EFFECT; ASSIGNMENT

This Agreement shall be binding upon and inure to the benefit of the parties hereto and their respective successors and permitted assigns. This Agreement shall not be assignable by Customer without QIAGEN's prior written consent; except that Customer may assign this Agreement, without the prior consent of QIAGEN, to the successor to all or substantially all of the business of Customer as long as the successor or surviving entity in such transaction agrees to be bound, in writing, by the terms and provisions of this Agreement, and written notice of such assignment is provided to QIAGEN prior to consummation of the transaction. This Agreement shall be assignable by QIAGEN.

18. SEVERABILITY

If any part of this Agreement shall be invalid or unenforceable under applicable law, such part shall be ineffective only to the extent of such invalidity or unenforceability, without in any way affecting the remaining parts of this Agreement. In addition, the part that is ineffective shall be reformed in a mutually agreeable manner so as to as most closely approximate, to the extent possible, the intent of the parties hereto.

19. SURVIVAL

The provisions of this Agreement that may reasonably be interpreted or construed as surviving the expiration or termination of this Agreement (including, without limitation, confidentiality and governing law) shall so survive for the period specified, or if no such period, for the applicable statute of limitations.

20. INDEPENDENT CONTRACTORS

The parties hereto are independent contractors and nothing in this Agreement will constitute the parties to be partners, nor constitute one party the agent of the other party, nor constitute the relationship to be a joint venture. Neither party shall have, or shall represent that it has,

the authority or power to act for or to undertake or create any obligation or responsibility, express or implied, on behalf of, or in the name of the other party.

21. GOVERNING LAW

This Agreement and any claims, disputes or causes of action relating to or arising out of this Agreement shall be construed in accordance with and governed by the laws of Germany without giving effect to the conflict of laws principles thereof. Notwithstanding any other provision under this Agreement, the parties to this Agreement submit to the exclusive jurisdiction of the Düsseldorf courts.

22. FORCE MAJEURE

QIAGEN shall not be liable for failure of or delay in performing obligations set forth in this Agreement, and shall not be deemed in breach of its obligations, if such failure or delay results from any of the following: Civil disobedience, hostilities, sabotage, terrorism, military actions, expropriation, nationalization or the escalation of any of the foregoing, any hurricane, flood, tornado, earthquake or other natural disaster, changes in weather conditions, epidemic, plague, pandemic or any other outbreak of illness, any law or regulation or any action taken by a government or public authority, including but not limited to an export or import restriction or other public health event in any country or any other event or circumstance outside of QIAGEN's reasonable control (each a "Force Majeure Event"). In such Force Majeure Event QIAGEN shall (a) promptly notify the Customer in writing and (b) use commercially reasonable efforts to cure or overcome the same and resume performance of its obligations hereunder. If such Force Majeure Event shall continue for a period of more than one calendar month, QIAGEN may terminate this Agreement without liability upon written notice to the Customer.

23. FINAL PROVISIONS

- a. Audits and inspections. Representatives of the Customer shall be entitled at reasonable times and with reasonable frequency during the QIAGEN's regular business hours,

and with reasonable advance notice (provided that no such advance notice shall be required for audits, which are required, by the applicable regulatory authority, to be conducted without prior notice), to visit the QIAGEN Genomic Services Sites in order to audit its facilities, equipment and procedures used for providing the Services for quality assurance purposes and to observe the progress of the Services. The auditing will take place no more than once per year and will last no longer than 1 working day Monday till Friday between 8:00 and 18:00. The provision of information, as well as audits or inspections referred to above are carried out at the expense and risk of the Customer.

- b. No amendment or modification of the terms of this Agreement shall be binding on either party unless in writing and signed on behalf of each party.