**Member Rights & Responsibilities**

1. Members have a right to receive information about the organization, its services, its network clinicians, and members’ rights and responsibilities.
2. Members have a right to be treated with respect and recognition of their dignity and right to privacy.
3. Members have a right to participate with network clinicians in making decisions about their health care.
4. Members have a right to a candid discussion of appropriate or medically necessary treatment options for their conditions, regardless of cost or benefit coverage.
5. Members have a right to voice complaints or appeals about the organization or the services it provides.
6. Members have a right to make recommendations regarding the organization’s members’ rights and responsibilities policies.
7. Members have a right to supply information (to the extent possible) that the organization and its network clinicians need in order to provide care.
8. Members have a right to a candid discussion of appropriate or medically necessary treatment options for their conditions.
9. Members have a right to be informed of their rights and responsibilities in a language they understand.
10. Members have a right to reasonable access to care, regardless of race, religion, gender, sexual orientation, ethnicity, age, or disability.
11. Members have a right to personal privacy and confidentiality of information.
12. Members have a right to have utilization management decisions made based on appropriateness of care.
13. Members have a right to be informed of the reason for any adverse determination, including the specific utilization review criteria or benefit provisions used in the determination.
14. Members have a right to designation of a surrogate decision-maker if the member is incapable of understanding a proposed treatment or procedure or is unable to communicate his or her wishes regarding care.
15. Members have a right to be informed of their rights and responsibilities in a language they understand.
16. If a member chooses not to comply with recommended care, treatment, or procedures, the clinician is to inform the member of the potential consequences of not complying with the treatment recommendations.
17. Members have a right to have utilization management decisions made based on appropriateness of care. The organization does not reward network clinicians or other individuals conducting utilization review for issuing adverse determinations for coverage or service.
18. Members have a right to be informed of rules and regulations concerning their own conduct.
19. Members have a right to reasonable access to care, regardless of race, religion, gender, sexual orientation, ethnicity, age, or disability.
20. Members have a right to be informed of the reason for any adverse determination, including the specific utilization review criteria or benefit provisions used in the determination.
21. Members have a right to have utilization management decisions made based on appropriateness of care. The organization does not reward network clinicians or other individuals conducting utilization review for issuing adverse determinations for coverage or service.
22. Members have a right to:
   - Inspect and copy their protected health information (PHI).
   - Request to amend their PHI.
   - Request an accounting of non-routine disclosures of PHI.
   - Request limitations on the use or disclosure of PHI.
   - Request confidential communications of PHI to be sent to an alternate address or by alternate means.
   - Make a complaint regarding use or disclosure of PHI.
   - Receive a Privacy Notice.
23. Members have a right to receive information about the organization’s clinical guidelines and Quality Improvement program.

The “Client Rights” shown in bold type are currently not being met by Providers and Payers in Idaho.