Canadian Code of Ethics for Psychologists

Third Edition
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Canadian Psychological Association
Société canadienne de psychologie

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Preamble

Introduction

Every discipline that has relatively autonomous control over its entry requirements, training, development of knowledge, standards, methods, and practices does so only within the context of a contract with the society in which it functions. This social contract is based on attitudes of mutual respect and trust, with society granting support for the autonomy of a discipline in exchange for a commitment by the discipline to do everything it can to assure that its members act ethically in conducting the affairs of the discipline within society; in particular, a commitment to try to assure that each member will place the welfare of the society and individual members of that society above the welfare of the discipline and its own members. By virtue of this social contract, psychologists have a higher duty of care to members of society than the general duty of care that all members of society have to each other.

The Canadian Psychological Association recognizes its responsibility to help assure ethical behaviour and attitudes on the part of psychologists. Attempts to assure ethical behaviour and attitudes include articulating ethical principles, values, and standards; promoting those principles, values, and standards through education, peer modelling, and consultation; developing and implementing methods to help psychologists monitor the ethics of their behaviour and attitudes; adjudicating complaints of unethical behaviour; and, taking corrective action when warranted.

This Code articulates ethical principles, values, and standards to guide all members of the Canadian Psychological Association, whether scientists, practitioners, or scientist practitioners, or whether acting in a research, direct service, teaching, student, trainee, administrative, management, employer, employee, supervisory, consultative, peer review, editorial, expert witness, social policy, or any other role related to the discipline of psychology.

Structure and Derivation of Code

Structure. Four ethical principles, to be considered and balanced in ethical decision making, are presented. Each principle is followed by a statement of those values that are included in and give definition to the principle. Each values statement is followed by a list of ethical standards that illustrate the application of the specific principle and values to the activities of psychologists. The standards range from minimal behavioural expectations (e.g., Standards I.28, II.28, III.33, IV.27) to more idealized, but achievable, attitudinal and behavioural expectations (e.g., Standards I.12, II.12, III.10, IV.6). In the margin, to the left of the standards, key words are placed to guide the reader through the standards and to illustrate the relationship of the specific standards to the values statement.

Derivation. The four principles represent those ethical principles used most consistently by Canadian psychologists to resolve hypothetical ethical dilemmas sent to them by the CPA Committee on Ethics during the initial development of the Code. In addition to the responses provided by Canadian psychologists, the values statements and ethical standards have been derived from interdisciplinary and international ethics codes, provincial and specialty codes of conduct, and ethics literature.
When Principles Conflict

All four principles are to be taken into account and balanced in ethical decision making. However, there are circumstances in which ethical principles will conflict and it will not be possible to give each principle equal weight. The complexity of ethical conflicts precludes a firm ordering of the principles. However, the four principles have been ordered according to the weight each generally should be given when they conflict, namely:

**Principle I: Respect for the Dignity of Persons.** This principle, with its emphasis on moral rights, generally should be given the highest weight, except in circumstances in which there is a clear and imminent danger to the physical safety of any person.

**Principle II: Responsible Caring.** This principle generally should be given the second highest weight. Responsible caring requires competence and should be carried out only in ways that respect the dignity of persons.

**Principle III: Integrity in Relationships.** This principle generally should be given the third highest weight. Psychologists are expected to demonstrate the highest integrity in all of their relationships. However, in rare circumstances, values such as openness and straightforwardness might need to be subordinated to the values contained in the Principles of Respect for the Dignity of Persons and Responsible Caring.

**Principle IV: Responsibility to Society.** This principle generally should be given the lowest weight of the four principles when it conflicts with one or more of them. Although it is necessary and important to consider responsibility to society in every ethical decision, adherence to this principle must be subject to and guided by Respect for the Dignity of Persons, Responsible Caring, and Integrity in Relationships. When a person’s welfare appears to conflict with benefits to society, it is often possible to find ways of working for the benefit of society that do not violate respect and responsible caring for the person. However, if this is not possible, the dignity and well-being of a person should not be sacrificed to a vision of the greater good of society, and greater weight must be given to respect and responsible caring for the person.

Even with the above ordering of the principles, psychologists will be faced with ethical dilemmas that are difficult to resolve. In these circumstances, psychologists are expected to engage in an ethical decision-making process that is explicit enough to bear public scrutiny. In some cases, resolution might be a matter of personal conscience. However, decisions of personal conscience are also expected to be the result of a decision-making process that is based on a reasonably coherent set of ethical principles and that can bear public scrutiny. If the psychologist can demonstrate that every reasonable effort was made to apply the ethical principles of this Code and resolution of the conflict has had to depend on the personal conscience of the psychologist, such a psychologist would be deemed to have followed this Code.

**The Ethical Decision-Making Process**

The ethical decision-making process might occur very rapidly, leading to an easy resolution of an ethical issue. This is particularly true of issues for which clear-cut guidelines or standards exist and for which there is no conflict between principles. On the other hand, some ethical issues (particularly those in which ethical principles conflict) are not easily resolved, might be emotionally distressful, and might require time-consuming deliberation.
The following basic steps typify approaches to ethical decision making:

1. Identification of the individuals and groups potentially affected by the decision.

2. Identification of ethically relevant issues and practices, including the interests, rights, and any relevant characteristics of the individuals and groups involved and of the system or circumstances in which the ethical problem arose.

3. Consideration of how personal biases, stresses, or self-interest might influence the development of or choice between courses of action.


5. Analysis of likely short-term, ongoing, and long-term risks and benefits of each course of action on the individual(s)/group(s) involved or likely to be affected (e.g., client, client’s family or employees, employing institution, students, research participants, colleagues, the discipline, society, self).

6. Choice of course of action after conscientious application of existing principles, values, and standards.

7. Action, with a commitment to assume responsibility for the consequences of the action.

8. Evaluation of the results of the course of action.

9. Assumption of responsibility for consequences of action, including correction of negative consequences, if any, or re-engaging in the decision-making process if the ethical issue is not resolved.

10. Appropriate action, as warranted and feasible, to prevent future occurrences of the dilemma (e.g., communication and problem solving with colleagues; changes in procedures and practices).

Psychologists engaged in time-consuming deliberation are encouraged and expected to consult with parties affected by the ethical problem, when appropriate, and with colleagues and/or advisory bodies when such persons can add knowledge or objectivity to the decision-making process. Although the decision for action remains with the individual psychologist, the seeking and consideration of such assistance reflects an ethical approach to ethical decision making.

Uses of the Code

This Code is intended to guide psychologists in their everyday conduct, thinking, and planning, and in the resolution of ethical dilemmas; that is, it advocates the practice of both proactive and reactive ethics.

The Code also is intended to serve as an umbrella document for the development of codes of conduct or other more specific codes. For example, the Code could be used as an ethical framework for the identification of behaviours that would be considered enforceable in a jurisdiction, the violation of which would constitute misconduct; or, jurisdictions could identify those standards in the Code that would be considered of a more serious nature and, therefore, reportable and subject to possible
discipline. In addition, the principles and values could be used to help specialty areas develop standards that are specific to those areas. Some work in this direction has already occurred within CPA (e.g., Guidelines for the Use of Animals in Research and Instruction in Psychology, Guidelines for Non-Discriminatory Practice, Guidelines for Psychologists in Addressing Recovered Memories). The principles and values incorporated into this Code, insofar as they come to be reflected in other documents guiding the behaviour of psychologists, will reduce inconsistency and conflict between documents.

A third use of the Code is to assist in the adjudication of complaints against psychologists. A body charged with this responsibility is required to investigate allegations, judge whether unacceptable behaviour has occurred, and determine what corrective action should be taken. In judging whether unacceptable conduct has occurred, many jurisdictions refer to a code of conduct. Some complaints, however, are about conduct that is not addressed directly in a code of conduct. The Code provides an ethical framework for determining whether the complaint is of enough concern, either at the level of the individual psychologist or at the level of the profession as a whole, to warrant corrective action (e.g., discipline of the individual psychologist, general educational activities for members, or incorporation into the code of conduct). In determining corrective action for an individual psychologist, one of the judgments the adjudicating body needs to make is whether an individual conscientiously engaged in an ethical decision-making process and acted in good faith, or whether there was a negligent or willful disregard of ethical principles. The articulation of the ethical decision-making process contained in this Code provides guidance for making such judgements.

Responsibility of the Individual Psychologist

The discipline’s contract with society commits the discipline and its members to act as a moral community that develops its ethical awareness and sensitivity, educates new members in the ethics of the discipline, manages its affairs and its members in an ethical manner, is as self-correcting as possible, and is accountable both internally and externally.

However, responsibility for ethical action depends foremost on the integrity of each individual psychologist; that is, on each psychologist’s commitment to behave as ethically as possible in every situation. Acceptance to membership in the Canadian Psychological Association, a scientific and professional association of psychologists, commits members:

1. To adhere to the Association’s Code in all current activities as a psychologist.

2. To apply conscientiously the ethical principles and values of the Code to new and emerging areas of activity.

3. To assess and discuss ethical issues and practices with colleagues on a regular basis.

4. To bring to the attention of the Association ethical issues that require clarification or the development of new guidelines or standards.

5. To bring concerns about possible unethical actions by a psychologist directly to the psychologist when the action appears to be primarily a lack of sensitivity, knowledge, or experience, and attempt to reach an agreement on the issue and, if needed, on the appropriate action to be taken.

6. To bring concerns about possible unethical actions of a more serious nature (e.g., actions that have caused or could cause serious harm, or actions that are considered
misconduct in the jurisdiction) to the person(s) or body(ies) best suited to investigating
the situation and to stopping or offsetting the harm.

7. To consider seriously others’ concerns about one’s own possibly unethical actions and
attempt to reach an agreement on the issue and, if needed, take appropriate action.

8. In bringing or in responding to concerns about possible unethical actions, not to be
vexatious or malicious.

9. To cooperate with duly constituted committees of the Association that are concerned
with ethics and ethical conduct.

Relationship of Code to Personal Behaviour

This Code is intended to guide and regulate only those activities a psychologist engages in by virtue of
being a psychologist. There is no intention to guide or regulate a psychologist’s activities outside of this
context. Personal behaviour becomes a concern of the discipline only if it is of such a nature that it
undermines public trust in the discipline as a whole or if it raises questions about the psychologist’s
ability to carry out appropriately his/her responsibilities as a psychologist.

Relationship of Code to Provincial Regulatory Bodies

In exercising its responsibility to articulate ethical principles, values, and standards for those who wish
to become and remain members in good standing, the Canadian Psychological Association recognizes
the multiple memberships that some psychologists have (both regulatory and voluntary). The Code has
attempted to encompass and incorporate those ethical principles most prevalent in the discipline as a
whole, thereby minimizing the possibility of variance with provincial/territorial regulations and
guidelines. Psychologists are expected to respect the requirements of their provincial/territorial
regulatory bodies. Such requirements might define particular behaviours that constitute misconduct, are
reportable to the regulatory body, and/or are subject to discipline.

Definition of Terms

For the purposes of this Code:

a) “Psychologist” means any person who is a Fellow, Member, Student Affiliate
or Foreign Affiliate of the Canadian Psychological Association, or a member
of any psychology voluntary association or regulatory body adopting this
Code. (Readers are reminded that provincial/territorial jurisdictions might
restrict the legal use of the term psychologist in their jurisdiction and that such
restrictions are to be honoured.)

b) “Client” means an individual, family, or group (including an organization or
community) receiving service from a psychologist.

c) Clients, research participants, students, and any other persons with whom
psychologists come in contact in the course of their work, are “independent”
if they can independently contract or give informed consent. Such persons are
“partially dependent” if the decision to contract or give informed consent is
shared between two or more parties (e.g., parents and school boards, workers
and Workers’ Compensation Boards, adult members of a family). Such

persons are considered to be “fully dependent” if they have little or no choice about whether or not to receive service or participate in an activity (e.g., patients who have been involuntarily committed to a psychiatric facility, or very young children involved in a research project).

d) “Others” means any persons with whom psychologists come in contact in the course of their work. This may include, but is not limited to: clients seeking help with individual, family, organizational, industrial, or community issues; research participants; employees; students; trainees; supervisees; colleagues; employers; third party payers; and, members of the general public.

e) “Legal or civil rights” means those rights protected under laws and statutes recognized by the province or territory in which the psychologist is working.

f) “Moral rights” means fundamental and inalienable human rights that might or might not be fully protected by existing laws and statutes. Of particular significance to psychologists, for example, are rights to: distributive justice; fairness and due process; and, developmentally appropriate privacy, self-determination, and personal liberty. Protection of some aspects of these rights might involve practices that are not contained or controlled within current laws and statutes. Moral rights are not limited to those mentioned in this definition.

g) “Unjust discrimination” or “unjustly discriminatory” means activities that are prejudicial or promote prejudice to persons because of their culture, nationality, ethnicity, colour, race, religion, sex, gender, marital status, sexual orientation, physical or mental abilities, age, socio-economic status, or any other preference or personal characteristic, condition, or status.

h) “Sexual harassment” includes either or both of the following: (i) The use of power or authority in an attempt to coerce another person to engage in or tolerate sexual activity. Such uses include explicit or implicit threats of reprisal for noncompliance, or promises of reward for compliance. (ii) Engaging in deliberate and/or repeated unsolicited sexually oriented comments, anecdotes, gestures, or touching, if such behaviours: are offensive and unwelcome; create an offensive, hostile, or intimidating working, learning, or service environment; or, can be expected to be harmful to the recipient.

i) The “discipline of psychology” refers to the scientific and applied methods and knowledge of psychology, and to the structures and procedures used by its members for conducting their work in relationship to society, to members of the public, to students or trainees, and to each other.

Review Schedule

To maintain the relevance and responsiveness of this Code, it will be reviewed regularly by the CPA Board of Directors, and revised as needed. You are invited to forward comments and suggestions, at any time, to the CPA office. In addition to psychologists, this invitation is extended to all readers, including members of the public and other disciplines.

Principle I: Respect for the Dignity of Persons

Values Statement

In the course of their work as scientists, practitioners, or scientist-practitioners, psychologists come into contact with many different individuals and groups, including: research participants; clients seeking help with individual, family, organizational, industrial, or community issues; students; trainees; supervisees; employees; business partners; business competitors; colleagues; employers; third party payers; and, the general public.

In these contacts, psychologists accept as fundamental the principle of respect for the dignity of persons; that is, the belief that each person should be treated primarily as a person or an end in him/herself, not as an object or a means to an end. In so doing, psychologists acknowledge that all persons have a right to have their innate worth as human beings appreciated and that this worth is not dependent upon their culture, nationality, ethnicity, colour, race, religion, sex, gender, marital status, sexual orientation, physical or mental abilities, age, socio-economic status, or any other preference or personal characteristic, condition, or status.

Although psychologists have a responsibility to respect the dignity of all persons with whom they come in contact in their role as psychologists, the nature of their contract with society demands that their greatest responsibility be to those persons in the most vulnerable position. Normally, persons directly receiving or involved in the psychologist’s activities are in such a position (e.g., research participants, clients, students). This responsibility is almost always greater than their responsibility to those indirectly involved (e.g., employers, third party payers, the general public).

Adherence to the concept of moral rights is an essential component of respect for the dignity of persons. Rights to privacy, self-determination, personal liberty, and natural justice are of particular importance to psychologists, and they have a responsibility to protect and promote these rights in all of their activities. As such, psychologists have a responsibility to develop and follow procedures for informed consent, confidentiality, fair treatment, and due process that are consistent with those rights.

As individual rights exist within the context of the rights of others and of responsible caring (see Principle II), there might be circumstances in which the possibility of serious detrimental consequences to themselves or others, a diminished capacity to be autonomous, or a court order, would disallow some aspects of the rights to privacy, self-determination, and personal liberty. Indeed, such circumstances might be serious enough to create a duty to warn or protect others (see Standards I.45 and II.39). However, psychologists still have a responsibility to respect the rights of the person(s) involved to the greatest extent possible under the circumstances, and to do what is necessary and reasonable to reduce the need for future disallowances.

Psychologists recognize that, although all persons possess moral rights, the manner in which such rights are promoted, protected, and exercised varies across communities and cultures. For instance, definitions of what is considered private vary, as does the role of families and other community members in personal decision making. In their work, psychologists acknowledge and respect such differences, while guarding against clear violations of moral rights.

In addition, psychologists recognize that as individual, family, group, or community vulnerabilities increase, or as the power of persons to control their environment or their lives decreases, psychologists have an increasing responsibility to seek ethical advice and to establish safeguards to protect the rights of the persons involved. For this reason, psychologists consider it their responsibility to increase safeguards to protect and promote the rights of persons involved in their activities proportionate to the
degree of dependency and the lack of voluntary initiation. For example, this would mean that there would be more safeguards to protect and promote the rights of fully dependent persons than partially dependent persons, and more safeguards for partially dependent than independent persons.

Respect for the dignity of persons also includes the concept of distributive justice. With respect to psychologists, this concept implies that all persons are entitled to benefit equally from the contributions of psychology and to equal quality in the processes, procedures, and services being conducted by psychologists, regardless of the person’s characteristics, condition, or status. Although individual psychologists might specialize and direct their activities to particular populations, or might decline to engage in activities based on the limits of their competence or acknowledgment of problems in some relationships, psychologists must not exclude persons on a capricious or unjustly discriminatory basis.

By virtue of the social contract that the discipline has with society, psychologists have a higher duty of care to members of society than the general duty of care all members of society have to each other. However, psychologists are entitled to protect themselves from serious violations of their own moral rights (e.g., privacy, personal liberty) in carrying out their work as psychologists.

**Ethical Standards**

In adhering to the Principle of Respect for the Dignity of Persons, psychologists would:

<p>| General respect | I.1 | Demonstrate appropriate respect for the knowledge, insight, experience, and areas of expertise of others. |
| General rights  | I.2 | Not engage publicly (e.g., in public statements, presentations, research reports, or with clients) in degrading comments about others, including demeaning jokes based on such characteristics as culture, nationality, ethnicity, colour, race, religion, sex, gender, or sexual orientation. |
|                | I.3 | Strive to use language that conveys respect for the dignity of persons as much as possible in all written or oral communication. |
|                | I.4 | Abstain from all forms of harassment, including sexual harassment. |
|                | I.5 | Avoid or refuse to participate in practices disrespectful of the legal, civil, or moral rights of others. |
|                | I.6 | Refuse to advise, train, or supply information to anyone who, in the psychologist’s judgement, will use the knowledge or skills to infringe on human rights. |
|                | I.7 | Make every reasonable effort to ensure that psychological knowledge is not misused, intentionally or unintentionally, to infringe on human rights. |
|                | I.8 | Respect the right of research participants, clients, employees, supervisees, students, trainees, and others to safeguard their own dignity. |</p>
<table>
<thead>
<tr>
<th>Non-discrimination</th>
<th>I.9</th>
<th>Not practice, condone, facilitate, or collaborate with any form of unjust discrimination.</th>
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<td></td>
<td>I.10</td>
<td>Act to correct practices that are unjustly discriminatory.</td>
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<td>I.11</td>
<td>Seek to design research, teaching, practice, and business activities in such a way that they contribute to the fair distribution of benefits to individuals and groups, and that they do not unfairly exclude those who are vulnerable or might be disadvantaged.</td>
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<td>Fair treatment/due process</td>
<td>I.12</td>
<td>Work and act in a spirit of fair treatment to others.</td>
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<td></td>
<td>I.13</td>
<td>Help to establish and abide by due process or other natural justice procedures for employment, evaluation, adjudication, editorial, and peer review activities.</td>
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<td>I.14</td>
<td>Compensate others fairly for the use of their time, energy, and knowledge, unless such compensation is refused in advance.</td>
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<td>I.15</td>
<td>Establish fees that are fair in light of the time, energy, and knowledge of the psychologist and any associates or employees, and in light of the market value of the product or service. (Also see Standard IV.12.)</td>
</tr>
<tr>
<td>Informed consent</td>
<td>I.16</td>
<td>Seek as full and active participation as possible from others in decisions that affect them, respecting and integrating as much as possible their opinions and wishes.</td>
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<td>I.17</td>
<td>Recognize that informed consent is the result of a process of reaching an agreement to work collaboratively, rather than of simply having a consent form signed.</td>
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<td>I.18</td>
<td>Respect the expressed wishes of persons to involve others (e.g., family members, community members) in their decision making regarding informed consent. This would include respect for written and clearly expressed unwritten advance directives.</td>
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<td></td>
<td>I.19</td>
<td>Obtain informed consent from all independent and partially dependent persons for any psychological services provided to them except in circumstances of urgent need (e.g., disaster or other crisis). In urgent circumstances, psychologists would proceed with the assent of such persons, but fully informed consent would be obtained as soon as possible. (Also see Standard I.29.)</td>
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<td>I.20</td>
<td>Obtain informed consent for all research activities that involve obtrusive measures, invasion of privacy, more than minimal risk of harm, or any attempt to change the behaviour of research participants.</td>
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<td></td>
<td>I.21</td>
<td>Establish and use signed consent forms that specify the dimensions of informed consent or that acknowledge that such dimensions</td>
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</table>
have been explained and are understood, if such forms are required by law or if such forms are desired by the psychologist, the person(s) giving consent, or the organization for whom the psychologist works.

I.22 Accept and document oral consent, in situations in which signed consent forms are not acceptable culturally or in which there are other good reasons for not using them.

I.23 Provide, in obtaining informed consent, as much information as reasonable or prudent persons would want to know before making a decision or consenting to the activity. The psychologist would relay this information in language that the persons understand (including providing translation into another language, if necessary) and would take whatever reasonable steps are needed to ensure that the information was, in fact, understood.

I.24 Ensure, in the process of obtaining informed consent, that at least the following points are understood: purpose and nature of the activity; mutual responsibilities; confidentiality protections and limitations; likely benefits and risks; alternatives; the likely consequences of non-action; the option to refuse or withdraw at any time, without prejudice; over what period of time the consent applies; and, how to rescind consent if desired. (Also see Standards III.23-30.)

I.25 Provide new information in a timely manner, whenever such information becomes available and is significant enough that it reasonably could be seen as relevant to the original or ongoing informed consent.

I.26 Clarify the nature of multiple relationships to all concerned parties before obtaining consent, if providing services to or conducting research at the request or for the use of third parties. This would include, but not be limited to: the purpose of the service or research; the reasonably anticipated use that will be made of information collected; and, the limits on confidentiality. Third parties may include schools, courts, government agencies, insurance companies, police, and special funding bodies.

Freedom of consent

I.27 Take all reasonable steps to ensure that consent is not given under conditions of coercion, undue pressure, or undue reward. (Also see Standard III.32.)

I.28 Not proceed with any research activity, if consent is given under any condition of coercion, undue pressure, or undue reward. (Also see Standard III.32.)

I.29 Take all reasonable steps to confirm or re-establish freedom of consent, if consent for service is given under conditions of duress or conditions of extreme need.
<table>
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<th>Paragraph</th>
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<tr>
<td>I.30</td>
<td>Respect the right of persons to discontinue participation or service at any time, and be responsive to non-verbal indications of a desire to discontinue if a person has difficulty with verbally communicating such a desire (e.g., young children, verbally disabled persons) or, due to culture, is unlikely to communicate such a desire orally.</td>
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<tr>
<td>I.31</td>
<td>Seek an independent and adequate ethical review of human rights issues and protections for any research involving members of vulnerable groups, including persons of diminished capacity to give informed consent, before making a decision to proceed.</td>
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<tr>
<td>I.32</td>
<td>Not use persons of diminished capacity to give informed consent in research studies, if the research involved may be carried out equally well with persons who have a fuller capacity to give informed consent.</td>
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<td>I.33</td>
<td>Seek to use methods that maximize the understanding and ability to consent of persons of diminished capacity to give informed consent, and that reduce the need for a substitute decision maker.</td>
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<td>I.34</td>
<td>Carry out informed consent processes with those persons who are legally responsible or appointed to give informed consent on behalf of persons not competent to consent on their own behalf, seeking to ensure respect for any previously expressed preferences of persons not competent to consent.</td>
</tr>
<tr>
<td>I.35</td>
<td>Seek willing and adequately informed participation from any person of diminished capacity to give informed consent, and proceed without this assent only if the service or research activity is considered to be of direct benefit to that person.</td>
</tr>
<tr>
<td>I.36</td>
<td>Be particularly cautious in establishing the freedom of consent of any person who is in a dependent relationship to the psychologist (e.g., student, employee). This may include, but is not limited to, offering that person an alternative activity to fulfill their educational or employment goals, or offering a range of research studies or experience opportunities from which the person can select, none of which is so onerous as to be coercive.</td>
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<tr>
<td>I.37</td>
<td>Seek and collect only information that is germane to the purpose(s) for which consent has been obtained.</td>
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<tr>
<td>I.38</td>
<td>Take care not to infringe, in research, teaching, or service activities, on the personally, developmentally, or culturally defined private space of individuals or groups, unless clear permission is granted to do so.</td>
</tr>
</tbody>
</table>
I.39 Record only that private information necessary for the provision of continuous, coordinated service, or for the goals of the particular research study being conducted, or that is required or justified by law. (Also see Standards IV.17 and IV.18.)

I.40 Respect the right of research participants, employees, supervisees, students, and trainees to reasonable personal privacy.

I.41 Collect, store, handle, and transfer all private information, whether written or unwritten (e.g., communication during service provision, written records, e-mail or fax communication, computer files, video-tapes), in a way that attends to the needs for privacy and security. This would include having adequate plans for records in circumstances of one’s own serious illness, termination of employment, or death.

I.42 Take all reasonable steps to ensure that records over which they have control remain personally identifiable only as long as necessary in the interests of those to whom they refer and/or to the research project for which they were collected, or as required or justified by law (e.g., the possible need to defend oneself against future allegations), and render anonymous or destroy any records under their control that no longer need to be personally identifiable. (Also see Standards IV.17 and IV.18.)

Confidentiality

I.43 Be careful not to relay information about colleagues, colleagues’ clients, research participants, employees, supervisees, students, trainees, and members of organizations, gained in the process of their activities as psychologists, that the psychologist has reason to believe is considered confidential by those persons, except as required or justified by law. (Also see Standards IV.17 and IV.18.)

I.44 Clarify what measures will be taken to protect confidentiality, and what responsibilities family, group, and community members have for the protection of each other’s confidentiality, when engaged in services to or research with individuals, families, groups, or communities.

I.45 Share confidential information with others only with the informed consent of those involved, or in a manner that the persons involved cannot be identified, except as required or justified by law, or in circumstances of actual or possible serious physical harm or death. (Also see Standards II.39, IV.17, and IV.18.)

Extended responsibility

I.46 Encourage others, in a manner consistent with this Code, to respect the dignity of persons and to expect respect for their own dignity.

I.47 Assume overall responsibility for the scientific and professional activities of their assistants, employees, students, supervisees, and trainees with regard to Respect for the Dignity of Persons, all of whom, however, incur similar obligations.
Principle II: Responsible Caring

Values Statement

A basic ethical expectation of any discipline is that its activities will benefit members of society or, at least, do no harm. Therefore, psychologists demonstrate an active concern for the welfare of any individual, family, group, or community with whom they relate in their role as psychologists. This concern includes both those directly involved and those indirectly involved in their activities. However, as with Principle I, psychologists’ greatest responsibility is to protect the welfare of those in the most vulnerable position. Normally, persons directly involved in their activities (e.g., research participants, clients, students) are in such a position. Psychologists’ responsibility to those indirectly involved (e.g., employers, third party payers, the general public) normally is secondary.

As persons usually consider their own welfare in their personal decision making, obtaining informed consent (see Principle I) is one of the best methods for ensuring that their welfare will be protected. However, it is only when such consent is combined with the responsible caring of the psychologist that there is considerable ethical protection of the welfare of the person(s) involved.

Responsible caring leads psychologists to take care to discern the potential harm and benefits involved, to predict the likelihood of their occurrence, to proceed only if the potential benefits outweigh the potential harms, to develop and use methods that will minimize harms and maximize benefits, and to take responsibility for correcting clearly harmful effects that have occurred as a direct result of their research, teaching, practice, or business activities.

In order to carry out these steps, psychologists recognize the need for competence and self-knowledge. They consider incompetent action to be unethical per se, as it is unlikely to be of benefit and likely to be harmful. They engage only in those activities in which they have competence or for which they are receiving supervision, and they perform their activities as competently as possible. They acquire, contribute to, and use the existing knowledge most relevant to the best interests of those concerned. They also engage in self-reflection regarding how their own values, attitudes, experiences, and social context (e.g., culture, ethnicity, colour, religion, sex, gender, sexual orientation, physical and mental abilities, age, and socio-economic status) influence their actions, interpretations, choices, and recommendations. This is done with the intent of increasing the probability that their activities will benefit and not harm the individuals, families, groups, and communities to whom they relate in their role as psychologists. Psychologists define harm and benefit in terms of both physical and psychological dimensions. They are concerned about such factors as: social, family, and community relationships; personal and cultural identity; feelings of self-worth, fear, humiliation, interpersonal trust, and cynicism; self-knowledge and general knowledge; and, such factors as physical safety, comfort, pain, and injury. They are concerned about immediate, short-term, and long-term effects.

Responsible caring recognizes and respects (e.g., through obtaining informed consent) the ability of individuals, families, groups, and communities to make decisions for themselves and to care for themselves and each other. It does not replace or undermine such ability, nor does it substitute one person’s opinion about what is in the best interests of another person for that other person’s competent decision making. However, psychologists recognize that, as vulnerabilities increase or as power to control one’s own life decreases, psychologists have an increasing responsibility to protect the well-being of the individual, family, group, or community involved. For this reason, as in Principle I, psychologists consider it their responsibility to increase safeguards proportionate to the degree of dependency and the lack of voluntary initiation on the part of the persons involved. However, for Principle II, the safeguards are for the well-being of persons rather than for the rights of persons.
Psychologists’ treatment and use of animals in their research and teaching activities are also a component of responsible caring. Although animals do not have the same moral rights as persons (e.g., privacy), they do have the right to be treated humanely and not to be exposed to unnecessary discomfort, pain, or disruption.

By virtue of the social contract that the discipline has with society, psychologists have a higher duty of care to members of society than the general duty of care all members of society have to each other. However, psychologists are entitled to protect their own basic well-being (e.g., physical safety, family relationships) in their work as psychologists.

**Ethical Standards**

In adhering to the Principle of Responsible Caring, psychologists would:

**General caring**

II.1 Protect and promote the welfare of clients, research participants, employees, supervisees, students, trainees, colleagues, and others.

II.2 Avoid doing harm to clients, research participants, employees, supervisees, students, trainees, colleagues, and others.

II.3 Accept responsibility for the consequences of their actions.

II.4 Refuse to advise, train, or supply information to anyone who, in the psychologist’s judgment, will use the knowledge or skills to harm others.

II.5 Make every reasonable effort to ensure that psychological knowledge is not misused, intentionally or unintentionally, to harm others.

**Competence and self-knowledge**

II.6 Offer or carry out (without supervision) only those activities for which they have established their competence to carry them out to the benefit of others.

II.7 Not delegate activities to persons not competent to carry them out to the benefit of others.

II.8 Take immediate steps to obtain consultation or to refer a client to a colleague or other appropriate professional, whichever is more likely to result in providing the client with competent service, if it becomes apparent that a client’s problems are beyond their competence.

II.9 Keep themselves up to date with a broad range of relevant knowledge, research methods, and techniques, and their impact on persons and society, through the reading of relevant literature, peer consultation, and continuing education activities, in order that their service or research activities and conclusions will benefit and not harm others.
II.10 Evaluate how their own experiences, attitudes, culture, beliefs, values, social context, individual differences, specific training, and stresses influence their interactions with others, and integrate this awareness into all efforts to benefit and not harm others.

II.11 Seek appropriate help and/or discontinue scientific or professional activity for an appropriate period of time, if a physical or psychological condition reduces their ability to benefit and not harm others.

II.12 Engage in self-care activities that help to avoid conditions (e.g., burnout, addictions) that could result in impaired judgment and interfere with their ability to benefit and not harm others.

Risk/benefit analysis

II.13 Assess the individuals, families, groups, and communities involved in their activities adequately enough to ensure that they will be able to discern what will benefit and not harm the persons involved.

II.14 Be sufficiently sensitive to and knowledgeable about individual, group, community, and cultural differences and vulnerabilities to discern what will benefit and not harm persons involved in their activities.

II.15 Carry out pilot studies to determine the effects of all new procedures and techniques that might carry more than minimal risk, before considering their use on a broader scale.

II.16 Seek an independent and adequate ethical review of the balance of risks and potential benefits of all research and new interventions that involve procedures of unknown consequence, or where pain, discomfort, or harm are possible, before making a decision to proceed.

II.17 Not carry out any scientific or professional activity unless the probable benefit is proportionately greater than the risk involved.

Maximize benefit

II.18 Provide services that are coordinated over time and with other service providers, in order to avoid duplication or working at cross purposes.

II.19 Create and maintain records relating to their activities that are sufficient to support continuity and appropriate coordination of their activities with the activities of others.

II.20 Make themselves aware of the knowledge and skills of other disciplines (e.g., law, medicine, business administration) and advise the use of such knowledge and skills, where relevant to the benefit of others.

II.21 Strive to provide and/or obtain the best possible service for those needing and seeking psychological service. This may include, but
is not limited to: selecting interventions that are relevant to the needs and characteristics of the client and that have reasonable theoretical or empirically-supported efficacy in light of those needs and characteristics; consulting with, or including in service delivery, persons relevant to the culture or belief systems of those served; advocating on behalf of the client; and, recommending professionals other than psychologists when appropriate.

II.22 Monitor and evaluate the effect of their activities, record their findings, and communicate new knowledge to relevant others.

II.23 Debrief research participants in such a way that the participants’ knowledge is enhanced and the participants have a sense of contribution to knowledge. (Also see Standards III.26 and III.27.)

II.24 Perform their teaching duties on the basis of careful preparation, so that their instruction is current and scholarly.

II.25 Facilitate the professional and scientific development of their employees, supervisees, students, and trainees by ensuring that these persons understand the values and ethical prescriptions of the discipline, and by providing or arranging for adequate working conditions, timely evaluations, and constructive consultation and experience opportunities.

II.26 Encourage and assist students in publication of worthy student papers.

Minimize harm

II.27 Be acutely aware of the power relationship in therapy and, therefore, not encourage or engage in sexual intimacy with therapy clients, neither during therapy, nor for that period of time following therapy during which the power relationship reasonably could be expected to influence the client’s personal decision making. (Also see Standard III.31.)

II.28 Not encourage or engage in sexual intimacy with students or trainees with whom the psychologist has an evaluative or other relationship of direct authority. (Also see Standard III.31.)

II.29 Be careful not to engage in activities in a way that could place incidentally involved persons at risk.

II.30 Be acutely aware of the need for discretion in the recording and communication of information, in order that the information not be misinterpreted or misused to the detriment of others. This includes, but is not limited to: not recording information that could lead to misinterpretation and misuse; avoiding conjecture; clearly labelling opinion; and, communicating information in language that can be understood clearly by the recipient of the information.
II.31 Give reasonable assistance to secure needed psychological services or activities, if personally unable to meet requests for needed psychological services or activities.

II.32 Provide a client, if appropriate and if desired by the client, with reasonable assistance to find a way to receive needed services in the event that third party payments are exhausted and the client cannot afford the fees involved.

II.33 Maintain appropriate contact, support, and responsibility for caring until a colleague or other professional begins service, if referring a client to a colleague or other professional.

II.34 Give reasonable notice and be reasonably assured that discontinuation will cause no harm to the client, before discontinuing services.

II.35 Screen appropriate research participants and select those least likely to be harmed, if more than minimal risk of harm to some research participants is possible.

II.36 Act to minimize the impact of their research activities on research participants’ personalities, or on their physical or mental integrity.

II.37 Terminate an activity when it is clear that the activity carries more than minimal risk of harm and is found to be more harmful than beneficial, or when the activity is no longer needed.

II.38 Refuse to help individuals, families, groups, or communities to carry out or submit to activities that, according to current knowledge, or legal or professional guidelines, would cause serious physical or psychological harm to themselves or others.

II.39 Do everything reasonably possible to stop or offset the consequences of actions by others when these actions are likely to cause serious physical harm or death. This may include reporting to appropriate authorities (e.g., the police), an intended victim, or a family member or other support person who can intervene, and would be done even when a confidential relationship is involved. (Also see Standard I.45.)

II.40 Act to stop or offset the consequences of seriously harmful activities being carried out by another psychologist or member of another discipline, when there is objective information about the activities and the harm, and when these activities have come to their attention outside of a confidential client relationship between themselves and the psychologist or member of another discipline. This may include reporting to the appropriate regulatory body, authority, or committee for action, depending on the psychologist’s judgment about the person(s) or body(ies) best suited to stop or
offset the harm, and depending upon regulatory requirements and definitions of misconduct.

II.41 Act also to stop or offset the consequences of harmful activities carried out by another psychologist or member of another discipline, when the harm is not serious or the activities appear to be primarily a lack of sensitivity, knowledge, or experience, and when the activities have come to their attention outside of a confidential client relationship between themselves and the psychologist or member of another discipline. This may include talking informally with the psychologist or member of the other discipline, obtaining objective information and, if possible and relevant, the assurance that the harm will discontinue and be corrected. If in a vulnerable position (e.g., employee, trainee) with respect to the other psychologist or member of the other discipline, it may include asking persons in less vulnerable positions to participate in the meeting(s).

II.42 Be open to the concerns of others about perceptions of harm that they as a psychologist might be causing, stop activities that are causing harm, and not punish or seek punishment for those who raise such concerns in good faith.

II.43 Not place an individual, group, family, or community needing service at a serious disadvantage by offering them no service in order to fulfill the conditions of a research design, when a standard service is available.

II.44 Debrief research participants in such a way that any harm caused can be discerned, and act to correct any resultant harm. (Also see Standards III.26 and III.27.)

Care of animals

II.45 Not use animals in their research unless there is a reasonable expectation that the research will increase understanding of the structures and processes underlying behaviour, or increase understanding of the particular animal species used in the study, or result eventually in benefits to the health and welfare of humans or other animals.

II.46 Use a procedure subjecting animals to pain, stress, or privation only if an alternative procedure is unavailable and the goal is justified by its prospective scientific, educational, or applied value.

II.47 Make every effort to minimize the discomfort, illness, and pain of animals. This would include performing surgical procedures only under appropriate anaesthesia, using techniques to avoid infection and minimize pain during and after surgery and, if disposing of experimental animals is carried out at the termination of the study, doing so in a humane way.
II.48 Use animals in classroom demonstrations only if the instructional objectives cannot be achieved through the use of video-tapes, films, or other methods, and if the type of demonstration is warranted by the anticipated instructional gain.

**Extended responsibility**

II.49 Encourage others, in a manner consistent with this *Code*, to care responsibly.

II.50 Assume overall responsibility for the scientific and professional activities of their assistants, employees, supervisees, students, and trainees with regard to the Principle of Responsible Caring, all of whom, however, incur similar obligations.
Principle III: Integrity in Relationships

Values Statement

The relationships formed by psychologists in the course of their work embody explicit and implicit mutual expectations of integrity that are vital to the advancement of scientific knowledge and to the maintenance of public confidence in the discipline of psychology. These expectations include: accuracy and honesty; straightforwardness and openness; the maximization of objectivity and minimization of bias; and, avoidance of conflicts of interest. Psychologists have a responsibility to meet these expectations and to encourage reciprocity.

In addition to accuracy, honesty, and the obvious prohibitions of fraud or misrepresentation, meeting expectations of integrity is enhanced by self-knowledge and the use of critical analysis. Although it can be argued that science is value-free and impartial, scientists are not. Personal values and self-interest can affect the questions psychologists ask, how they ask those questions, what assumptions they make, their selection of methods, what they observe and what they fail to observe, and how they interpret their data.

Psychologists are not expected to be value-free or totally without self-interest in conducting their activities. However, they are expected to understand how their backgrounds, personal needs, and values interact with their activities, to be open and honest about the influence of such factors, and to be as objective and unbiased as possible under the circumstances.

The values of openness and straightforwardness exist within the context of Respect for the Dignity of Persons (Principle I) and Responsible Caring (Principle II). As such, there will be circumstances in which openness and straightforwardness will need to be tempered. Fully open and straightforward disclosure might not be needed or desired by others and, in some circumstances, might be a risk to their dignity or well-being, or considered culturally inappropriate. In such circumstances, however, psychologists have a responsibility to ensure that their decision not to be fully open or straightforward is justified by higher-order values and does not invalidate any informed consent procedures.

Of special concern to psychologists is the provision of incomplete disclosure when obtaining informed consent for research participation, or temporarily leading research participants to believe that a research project has a purpose other than its actual purpose. These actions sometimes occur in research where full disclosure would be likely to influence the responses of the research participants and thus invalidate the results. Although research that uses such techniques can lead to knowledge that is beneficial, such benefits must be weighed against the research participant’s right to self-determination and the importance of public and individual trust in psychology. Psychologists have a serious obligation to avoid as much as possible the use of such research procedures. They also have a serious obligation to consider the need for, the possible consequences of, and their responsibility to correct any resulting mistrust or other harmful effects from their use.

As public trust in the discipline of psychology includes trusting that psychologists will act in the best interests of members of the public, situations that present real or potential conflicts of interest are of concern to psychologists. Conflict-of-interest situations are those that can lead to distorted judgment and can motivate psychologists to act in ways that meet their own personal, political, financial, or business interests at the expense of the best interests of members of the public. Although avoidance of all conflicts of interest and potential exploitation of others is not possible, some are of such a high risk to protecting the interests of members of the public and to maintaining the trust of the public, that they are considered never acceptable (see Standard III.31). The risk level of other conflicts of interest (e.g., dual or multiple relationships) might be partially dependent on cultural factors and the specific type of
professional relationship (e.g., long-term psychotherapy vs. community development activities). It is the responsibility of psychologists to avoid dual or multiple relationships and other conflicts of interest when appropriate and possible. When such situations cannot be avoided or are inappropriate to avoid, psychologists have a responsibility to declare that they have a conflict of interest, to seek advice, and to establish safeguards to ensure that the best interests of members of the public are protected.

Integrity in relationships implies that psychologists, as a matter of honesty, have a responsibility to maintain competence in any specialty area for which they declare competence, whether or not they are currently practising in that area. It also requires that psychologists, in as much as they present themselves as members and representatives of a specific discipline, have a responsibility to actively rely on and be guided by that discipline and its guidelines and requirements.

Ethical Standards

In adhering to the Principle of Integrity in Relationships, psychologists would:

**Accuracy/honesty**

| III.1 | Not knowingly participate in, condone, or be associated with dishonesty, fraud, or misrepresentation. |

| III.2 | Accurately represent their own and their colleagues’ credentials, qualifications, education, experience, competence, and affiliations, in all spoken, written, or printed communications, being careful not to use descriptions or information that could be misinterpreted (e.g., citing membership in a voluntary association of psychologists as a testament of competence). |

| III.3 | Carefully protect their own and their colleagues’ credentials from being misrepresented by others, and act quickly to correct any such misrepresentation. |

| III.4 | Maintain competence in their declared area(s) of psychological competence, as well as in their current area(s) of activity. (Also see Standard II.9.) |

| III.5 | Accurately represent their own and their colleagues’ activities, functions, contributions, and likely or actual outcomes of their activities (including research results) in all spoken, written, or printed communication. This includes, but is not limited to: advertisements of services or products; course and workshop descriptions; academic grading requirements; and, research reports. |

| III.6 | Ensure that their own and their colleagues’ activities, functions, contributions, and likely or actual outcomes of their activities (including research results) are not misrepresented by others, and act quickly to correct any such misrepresentation. |

| III.7 | Take credit only for the work and ideas that they have actually done or generated, and give credit for work done or ideas contributed by others (including students), in proportion to their contribution. |
III.8 Acknowledge the limitations of their own and their colleagues’ knowledge, methods, findings, interventions, and views.

III.9 Not suppress disconfirming evidence of their own and their colleagues’ findings and views, acknowledging alternative hypotheses and explanations.

**Objectivity/lack of bias**

III.10 Evaluate how their personal experiences, attitudes, values, social context, individual differences, stresses, and specific training influence their activities and thinking, integrating this awareness into all attempts to be objective and unbiased in their research, service, and other activities.

III.11 Take care to communicate as completely and objectively as possible, and to clearly differentiate facts, opinions, theories, hypotheses, and ideas, when communicating knowledge, findings, and views.

III.12 Present instructional information accurately, avoiding bias in the selection and presentation of information, and publicly acknowledge any personal values or bias that influence the selection and presentation of information.

III.13 Act quickly to clarify any distortion by a sponsor, client, agency (e.g., news media), or other persons, of the findings of their research.

**Straightforwardness/openness**

III.14 Be clear and straightforward about all information needed to establish informed consent or any other valid written or unwritten agreement (for example: fees, including any limitations imposed by third-party payers; relevant business policies and practices; mutual concerns; mutual responsibilities; ethical responsibilities of psychologists; purpose and nature of the relationship, including research participation; alternatives; likely experiences; possible conflicts; possible outcomes; and, expectations for processing, using, and sharing any information generated).

III.15 Provide suitable information about the results of assessments, evaluations, or research findings to the persons involved, if appropriate and if asked. This information would be communicated in understandable language.

III.16 Fully explain reasons for their actions to persons who have been affected by their actions, if appropriate and if asked.

III.17 Honour all promises and commitments included in any written or verbal agreement, unless serious and unexpected circumstances (e.g., illness) intervene. If such circumstances occur, then the psychologist would make a full and honest explanation to other parties involved.
III.18 Make clear whether they are acting as private citizens, as members of specific organizations or groups, or as representatives of the discipline of psychology, when making statements or when involved in public activities.

III.19 Carry out, present, and discuss research in a way that is consistent with a commitment to honest, open inquiry, and to clear communication of any research aims, sponsorship, social context, personal values, or financial interests that might affect or appear to affect the research.

III.20 Submit their research, in some accurate form and within the limits of confidentiality, to persons with expertise in the research area, for their comments and evaluations, prior to publication or the preparation of any final report.

III.21 Encourage and not interfere with the free and open exchange of psychological knowledge and theory between themselves, their students, colleagues, and the public.

III.22 Make no attempt to conceal the status of a trainee and, if a trainee is providing direct client service, ensure that the client is informed of that fact.

Avoidance of incomplete disclosure

III.23 Not engage in incomplete disclosure, or in temporarily leading research participants to believe that a research project or some aspect of it has a different purpose, if there are alternative procedures available or if the negative effects cannot be predicted or offset.

III.24 Not engage in incomplete disclosure, or in temporarily leading research participants to believe that a research project or some aspect of it has a different purpose, if it would interfere with the person’s understanding of facts that clearly might influence a decision to give adequately informed consent (e.g., withholding information about the level of risk, discomfort, or inconvenience).

III.25 Use the minimum necessary incomplete disclosure or temporary leading of research participants to believe that a research project or some aspect of it has a different purpose, when such research procedures are used.

III.26 Debrief research participants as soon as possible after the participants’ involvement, if there has been incomplete disclosure or temporary leading of research participants to believe that a research project or some aspect of it has a different purpose.

III.27 Provide research participants, during such debriefing, with a clarification of the nature of the study, seek to remove any misconceptions that might have arisen, and seek to re-establish any trust that might have been lost, assuring the participants that the
research procedures were neither arbitrary nor capricious, but necessary for scientifically valid findings. (Also see Standards II.23 and II.44.)

III.28 Act to re-establish with research participants any trust that might have been lost due to the use of incomplete disclosure or temporarily leading research participants to believe that the research project or some aspect of it had a different purpose.

III.29 Give a research participant the option of removing his or her data, if the research participant expresses concern during the debriefing about the incomplete disclosure or the temporary leading of the research participant to believe that the research project or some aspect of it had a different purpose, and if removal of the data will not compromise the validity of the research design and hence diminish the ethical value of the participation of the other research participants.

III.30 Seek an independent and adequate ethical review of the risks to public or individual trust and of safeguards to protect such trust for any research that plans to provide incomplete disclosure or temporarily lead research participants to believe that the research project or some aspect of it has a different purpose, before making a decision to proceed.

Avoidance of conflict of interest

III.31 Not exploit any relationship established as a psychologist to further personal, political, or business interests at the expense of the best interests of their clients, research participants, students, employers, or others. This includes, but is not limited to: soliciting clients of one’s employing agency for private practice; taking advantage of trust or dependency to encourage or engage in sexual intimacies (e.g., with clients not included in Standard II.27, with clients’ partners or relatives, with students or trainees not included in Standard II.28, or with research participants); taking advantage of trust or dependency to frighten clients into receiving services; misappropriating students’ ideas, research or work; using the resources of one’s employing institution for purposes not agreed to; giving or receiving kickbacks or bonuses for referrals; seeking or accepting loans or investments from clients; and, prejudicing others against a colleague for reasons of personal gain.

III.32 Not offer rewards sufficient to motivate an individual or group to participate in an activity that has possible or known risks to themselves or others. (Also see Standards I.27, I.28, II.2, and II.49.)

III.33 Avoid dual or multiple relationships (e.g., with clients, research participants, employees, supervisees, students, or trainees) and other situations that might present a conflict of interest or that might reduce their ability to be objective and unbiased in their determinations of what might be in the best interests of others.
III.34 Manage dual or multiple relationships that are unavoidable due to cultural norms or other circumstances in such a manner that bias, lack of objectivity, and risk of exploitation are minimized. This might include obtaining ongoing supervision or consultation for the duration of the dual or multiple relationship, or involving a third party in obtaining consent (e.g., approaching a client or employee about becoming a research participant).

III.35 Inform all parties, if a real or potential conflict of interest arises, of the need to resolve the situation in a manner that is consistent with Respect for the Dignity of Persons (Principle I) and Responsible Caring (Principle II), and take all reasonable steps to resolve the issue in such a manner.

Reliance on the discipline

III.36 Familiarize themselves with their discipline’s rules and regulations, and abide by them, unless abiding by them would be seriously detrimental to the rights or welfare of others as demonstrated in the Principles of Respect for the Dignity of Persons or Responsible Caring. (See Standards IV.17 and IV.18 for guidelines regarding the resolution of such conflicts.)

III.37 Familiarize themselves with and demonstrate a commitment to maintaining the standards of their discipline.

III.38 Seek consultation from colleagues and/or appropriate groups and committees, and give due regard to their advice in arriving at a responsible decision, if faced with difficult situations.

Extended responsibility

III.39 Encourage others, in a manner consistent with this Code, to relate with integrity.

III.40 Assume overall responsibility for the scientific and professional activities of their assistants, employees, supervisees, students, and trainees with regard to the Principle of Integrity in Relationships, all of whom, however, incur similar obligations.
Principle IV: Responsibility to Society

Values Statement

Psychology functions as a discipline within the context of human society. Psychologists, both in their work and as private citizens, have responsibilities to the societies in which they live and work, such as the neighbourhood or city, and to the welfare of all human beings in those societies.

Two of the legitimate expectations of psychology as a science and a profession are that it will increase knowledge and that it will conduct its affairs in such ways that it will promote the welfare of all human beings.

Freedom of enquiry and debate (including scientific and academic freedom) is a foundation of psychological education, science, and practice. In the context of society, the above expectations imply that psychologists will exercise this freedom through the use of activities and methods that are consistent with ethical requirements.

The above expectations also imply that psychologists will do whatever they can to ensure that psychological knowledge, when used in the development of social structures and policies, will be used for beneficial purposes, and that the discipline’s own structures and policies will support those beneficial purposes. Within the context of this document, social structures and policies that have beneficial purposes are defined as those that more readily support and reflect respect for the dignity of persons, responsible caring, integrity in relationships, and responsibility to society. If psychological knowledge or structures are used against these purposes, psychologists have an ethical responsibility to try to draw attention to and correct the misuse. Although this is a collective responsibility, those psychologists having direct involvement in the structures of the discipline, in social development, or in the theoretical or research data base that is being used (e.g., through research, expert testimony, or policy advice) have the greatest responsibility to act. Other psychologists must decide for themselves the most appropriate and beneficial use of their time and talents to help meet this collective responsibility.

In carrying out their work, psychologists acknowledge that many social structures have evolved slowly over time in response to human need and are valued by the societies that have developed them. In such circumstances, psychologists convey respect for such social structures and avoid unwarranted or unnecessary disruption. Suggestions for and action toward changes or enhancement of such structures are carried out through processes that seek to achieve a consensus within those societies and/or through democratic means.

On the other hand, if structures or policies seriously ignore or oppose the principles of respect for the dignity of persons, responsible caring, integrity in relationships, or responsibility to society, psychologists involved have a responsibility to speak out in a manner consistent with the principles of this Code, and advocate for appropriate change to occur as quickly as possible.

In order to be responsible and accountable to society, and to contribute constructively to its ongoing development, psychologists need to be willing to work in partnership with others, be self-reflective, and be open to external suggestions and criticisms about the place of the discipline of psychology in society. They need to engage in even-tempered observation and interpretation of the effects of societal structures and policies, and their process of change, developing the ability of psychologists to increase the

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1 Society is used here in the broad sense of a group of persons living as members of one or more human communities, rather than in the limited sense of state or government.
beneficial use of psychological knowledge and structures, and avoid their misuse. The discipline needs to be willing to set high standards for its members, to do what it can to assure that such standards are met, and to support its members in their attempts to maintain the standards. Once again, individual psychologists must decide for themselves the most appropriate and beneficial use of their time and talents in helping to meet these collective responsibilities.

**Ethical Standards**

In adhering to the Principle of Responsibility to Society, psychologists would:

<table>
<thead>
<tr>
<th>Development of knowledge</th>
<th>IV.1 Contribute to the discipline of psychology and of society’s understanding of itself and human beings generally, through free enquiry and the acquisition, transmission, and expression of knowledge and ideas, unless such activities conflict with other basic ethical requirements.</th>
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<td>IV.2 Not interfere with, or condone interference with, free enquiry and the acquisition, transmission, and expression of knowledge and ideas that do not conflict with other basic ethical requirements.</td>
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<td></td>
<td>IV.3 Keep informed of progress in their area(s) of psychological activity, take this progress into account in their work, and try to make their own contributions to this progress.</td>
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<tr>
<td>Beneficial activities</td>
<td>IV.4 Participate in and contribute to continuing education and the professional and scientific growth of self and colleagues.</td>
</tr>
<tr>
<td></td>
<td>IV.5 Assist in the development of those who enter the discipline of psychology by helping them to acquire a full understanding of their ethical responsibilities, and the needed competencies of their chosen area(s), including an understanding of critical analysis and of the variations, uses, and possible misuses of the scientific paradigm.</td>
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<td>IV.6 Participate in the process of critical self-evaluation of the discipline’s place in society, and in the development and implementation of structures and procedures that help the discipline to contribute to beneficial societal functioning and changes.</td>
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<td>IV.7 Provide and/or contribute to a work environment that supports the respectful expression of ethical concern or dissent, and the constructive resolution of such concern or dissent.</td>
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<td>IV.8 Engage in regular monitoring, assessment, and reporting (e.g., through peer review, and in programme reviews, case management reviews, and reports of one’s own research) of their ethical practices and safeguards.</td>
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<td>IV.9 Help develop, promote, and participate in accountability processes and procedures related to their work.</td>
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IV.10 Uphold the discipline’s responsibility to society by promoting and maintaining the highest standards of the discipline.

IV.11 Protect the skills, knowledge, and interpretations of psychology from being misused, used incompetently, or made useless (e.g., loss of security of assessment techniques) by others.

IV.12 Contribute to the general welfare of society (e.g., improving accessibility of services, regardless of ability to pay) and/or to the general welfare of their discipline, by offering a portion of their time to work for which they receive little or no financial return.

IV.13 Uphold the discipline’s responsibility to society by bringing incompetent or unethical behaviour, including misuses of psychological knowledge and techniques, to the attention of appropriate authorities, committees, or regulatory bodies, in a manner consistent with the ethical principles of this Code, if informal resolution or correction of the situation is not appropriate or possible.

IV.14 Enter only into agreements or contracts that allow them to act in accordance with the ethical principles and standards of this Code.

Respect for society

IV.15 Acquire an adequate knowledge of the culture, social structure, and customs of a community before beginning any major work there.

IV.16 Convey respect for and abide by prevailing community mores, social customs, and cultural expectations in their scientific and professional activities, provided that this does not contravene any of the ethical principles of this Code.

IV.17 Familiarize themselves with the laws and regulations of the societies in which they work, especially those that are related to their activities as psychologists, and abide by them. If those laws or regulations seriously conflict with the ethical principles contained herein, psychologists would do whatever they could to uphold the ethical principles. If upholding the ethical principles could result in serious personal consequences (e.g., jail or physical harm), decision for final action would be considered a matter of personal conscience.

IV.18 Consult with colleagues, if faced with an apparent conflict between abiding by a law or regulation and following an ethical principle, unless in an emergency, and seek consensus as to the most ethical course of action and the most responsible, knowledgeable, effective, and respectful way to carry it out.

Development of society

IV.19 Act to change those aspects of the discipline of psychology that detract from beneficial societal changes, where appropriate and possible.
IV.20 Be sensitive to the needs, current issues, and problems of society, when determining research questions to be asked, services to be developed, content to be taught, information to be collected, or appropriate interpretation of results or findings.

IV.21 Be especially careful to keep well informed of social issues through relevant reading, peer consultation, and continuing education, if their work is related to societal issues.

IV.22 Speak out, in a manner consistent with the four principles of this Code, if they possess expert knowledge that bears on important societal issues being studied or discussed.

IV.23 Provide thorough discussion of the limits of their data with respect to social policy, if their work touches on social policy and structure.

IV.24 Consult, if feasible and appropriate, with groups, organizations, or communities being studied, in order to increase the accuracy of interpretation of results and to minimize risk of misinterpretation or misuse.

IV.25 Make themselves aware of the current social and political climate and of previous and possible future societal misuses of psychological knowledge, and exercise due discretion in communicating psychological information (e.g., research results, theoretical knowledge), in order to discourage any further misuse.

IV.26 Exercise particular care when reporting the results of any work regarding vulnerable groups, ensuring that results are not likely to be misinterpreted or misused in the development of social policy, attitudes, and practices (e.g., encouraging manipulation of vulnerable persons or reinforcing discrimination against any specific population).

IV.27 Not contribute to nor engage in research or any other activity that contravenes international humanitarian law, such as the development of methods intended for use in the torture of persons, the development of prohibited weapons, or destruction of the environment.

IV.28 Provide the public with any psychological knowledge relevant to the public’s informed participation in the shaping of social policies and structures, if they possess expert knowledge that bears on the social policies and structures.

IV.29 Speak out and/or act, in a manner consistent with the four principles of this Code, if the policies, practices, laws, or regulations of the social structure within which they work seriously ignore or contradict any of the principles of this Code.
Extended responsibility

IV.30 Encourage others, in a manner consistent with this Code, to exercise responsibility to society.

IV.31 Assume overall responsibility for the scientific and professional activities of their assistants, employees, supervisees, students, and trainees with regard to the Principle of Responsibility to Society, all of whom, however, incur similar obligations.