Written directives for the administration of radiopharmaceuticals shall not be carried out if there are questions regarding the details of the administration.

To minimize the likelihood of a radiopharmaceutical misadministration.

1. Only those persons who are qualified by their training will carry out a written directive for the administration of a radiopharmaceutical.
2. If the person who is to perform a radiopharmaceutical administration receives conflicting information regarding patient identification, the administration is not to be performed until a positive identification can be made.
3. If the person who is to perform a radiopharmaceutical administration has some doubt about the appropriateness of the radiopharmaceutical, activity, or route of administration then guidance should be sought from the authorized user who signed the directive, or from another person trained in the performance of such a procedure rather than continuing the procedure when there is any doubt.
4. If the person who is to perform a radiopharmaceutical administration has a question about how the administration is to be performed, guidance should be sought from the authorized user who signed the directive, or from another person trained in the performance of such a procedure rather than continuing the procedure when there is any doubt.