



NEW MEXICO BEEF QUALITY ASSURANCE CERTIFICATION PROGRAM
Critical Management Plan / Affidavit of Compliance

Answer all questions as if you are responsible for the task or duty specified.

	Yes	No
1. Treatment procedures will comply with either label directions or as prescribed by a veterinarian with a valid veterinarian-client-patient relationship following FDA's policy for extra-label drug use.	<input type="checkbox"/>	<input type="checkbox"/>
2. All treatments administered extra-label will be kept to a minimum and will comply with the prescribed extended withdrawal time. Should there be any questions about withdrawal period compliance, a veterinarian will evaluate the treatment history against available withdrawal and residue information.	<input type="checkbox"/>	<input type="checkbox"/>
3. Mass medications and individual treatments given pre-weaning will be recorded on a group/pen basis. Individual treatment given post-weaning will be recorded on an individual basis. Records will consist of date, pen/individual identification, product used, amount given, location given, and withdrawal time.	<input type="checkbox"/>	<input type="checkbox"/>
4. A quality control program will be maintained for all incoming feed ingredients, including analyzing and eliminating suspect contamination such as molds, mycotoxins, pesticides and herbicides.	<input type="checkbox"/>	<input type="checkbox"/>
5. No banned ruminant-derived protein feedstuffs will be fed to cattle.	<input type="checkbox"/>	<input type="checkbox"/>
6. Only FDA approved medicated feed additives will be used, and only in accordance with label directions. No extra-label feed additives will be used.	<input type="checkbox"/>	<input type="checkbox"/>
7. Records of all medicated feed rations and batch runs will be maintained for at least two years by the feeders or managers who feed such rations.	<input type="checkbox"/>	<input type="checkbox"/>
8. All injections will be given subcutaneously in front of the shoulder. If products must be given intramuscularly, the product will be administered in the neck region, without exception. Products which cause tissue damage will be avoided when possible.	<input type="checkbox"/>	<input type="checkbox"/>
9. All injectable products administered will be limited to 10 cc per injection site.	<input type="checkbox"/>	<input type="checkbox"/>
10. The business will strive, during animal handling and transport, to prevent injuries and stresses that could lead to dark cutters and bruises.	<input type="checkbox"/>	<input type="checkbox"/>
11. Waterers, feed bunks, and other facilities will be managed to reduce pathogens and to minimize mud and manure on the hide of market cattle.	<input type="checkbox"/>	<input type="checkbox"/>
12. All cattle shipped to another facility or to slaughter will be checked to verify that withdrawal times have been met.	<input type="checkbox"/>	<input type="checkbox"/>
13. All records will be kept for two years. Appropriate New Mexico BQA/CMP documentation if requested will be transferred with the cattle as they move from one facility to another and will be available for inspection by the authorized auditing agent.	<input type="checkbox"/>	<input type="checkbox"/>

I understand the importance of the above listed items in the NM Beef Quality Assurance Program and agree to follow the recommended production practices.

Certification No. _____

Date _____

Printed Name _____

Email _____

Mailing Address _____

County _____

City, State, Zip _____

Phone _____

(check one) **Home** **Business** **Mobile**

Signature _____

In compliance with **BQA Recertification Guidelines**, this form must be completed and returned to the **County Extension Office/Agent**.