



# New Zealand Dairy Regulation Overview

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*Growing and Protecting New Zealand*



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# Summary of Presentation

1. Dairy legislation overview
2. Regulatory framework
3. Risk Management Programmes
4. Third party auditing, verification and laboratories
5. Monitoring by the regulator

# Regulation Overview

# Dairy Legislation Overview

<b>Primary Production</b>	<b>Primary Processing</b>	<b>Export</b>	<b>Domestic Sale</b>
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**Animal Products Act 1999**

**Ag Compounds and Vet  
Medicines Act 1997**

**Food Act 2014**

**Biosecurity Act 1993**

# Animal Products Act 1999

## Object of the Act

- Manage risks to human and animal health from animal material and products
- Ensure products are fit for the intended purpose
- Facilitate market access

## Coverage

- Primary processing of animal products

# Animal Products Act 1999

## Risk Management System

- Risk management programmes
- Regulated control schemes
- Official assurance safeguards
- Recognised agencies and persons
- Duties.

# Regulatory Structure

- **Animal Products Act**
  - Regulations
    - Standards, Specifications and Notices
      - Criteria
      - Codes of Practice
      - Templates (generic risk management plans)
      - Guidance

# Animal Products Regulations and Specifications

## Regulations

- Generic, outcome-oriented requirements that apply across industry  
eg Animal product standards, operating standards

## Specifications

- Provide technical detail or give effect to standards
- Are made by Director General, subject to consultation  
eg contaminant limits, premises hygiene, sampling and testing



# Animal Products Regulations and Specifications – Dairy Road Map

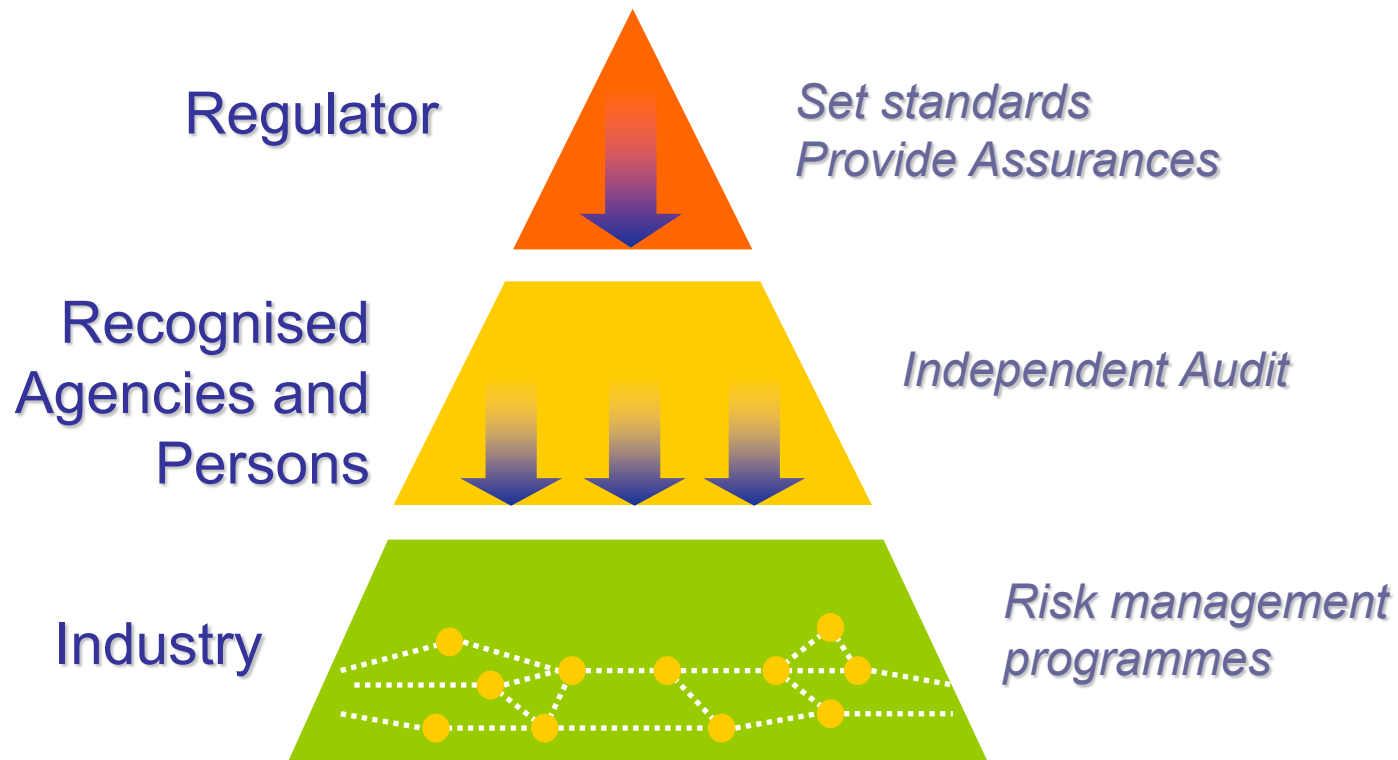
<http://www.foodsafety.govt.nz/elibrary/industry/dairy-legislation-roadmap.pdf>

# Regulatory Framework

# Roles and Responsibilities

- Ministry for Primary Industries (the Regulator)
- Third Party evaluators and verifiers of risk management programmes
- Farm dairy assessors
- Veterinarians
- Risk management programme operators
- Farm dairy operators (farmers)

# Regulatory Model



# Dairy Processing

# When an RMP is needed

## Dairy

- Farm dairies
- Dairy processors exporting

# Risk Management Programmes

- Documented programme
  - developed by company
- Manage known biological, chemical and physical hazards
- Wholesomeness
- Truthful labelling
- Products 'fit for intended purpose'.



# RMP Responsibilities





# What's In an RMP

**Scope**

**Authorities and Responsibilities**

**Product Description**

**Fitness for intended purpose**

**Process Description**

**Identification of risk factors:**

- Hazards (human, animals)
- Risks to wholesomeness,
- Risks from false or misleading labeling

**Control of risk factors**

**Provision for verification**

**(Processor's - unique requirements)**

**Documentation and records**

# Farm Dairies

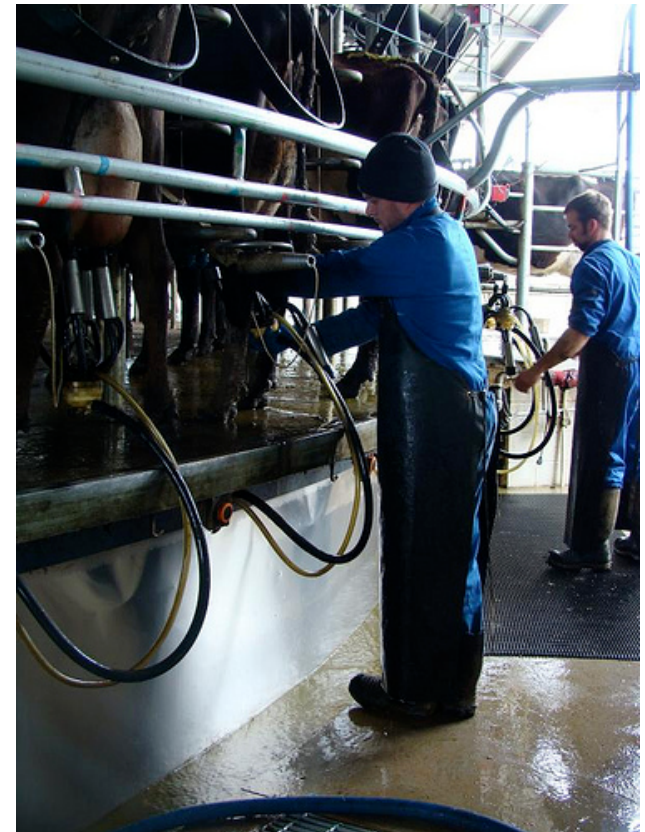
Risk management programmes for farm dairies must address:

- Siting, design and construction
- Equipment specifications
- Milk harvesting activities
- Animal health management
- Veterinary oversight
- Farm dairy water quality
- Milk cooling
- Control of chemicals
- Monitoring of milk quality



# Monitoring Farm Dairy Operations

- Farm Dairies must be independently assessed (audited) at least once a year by a Farm Dairy Assessor
- Farm Dairy Assessor competence is confirmed by a Recognised Verifier
- Farm Dairy Verifier competence is confirmed by accreditation bodies and MPI Compliance



# Raw Milk Acceptance

Raw milk acceptance requirements (applied at collection and delivery) include:

- Milk Cooling
- Sampling and testing
- Wholesomeness
- Microbiological parameters
- Residues and contaminants
- Animal health parameters



# Manufacturing

MPI sets the minimum outcome requirements for:

- Microbiological parameters (pathogens and process hygiene indicators)
- Chemical Residues and Contaminants
- Nutritional standards for specified foods (e.g. infant formula)
- Country specific requirements



# General Criteria

MPI sets minimum RMP requirements for:

- Reporting Requirements
- Dairy Product Safety Limits specified
- Residues of Agricultural Compounds & Veterinary Medicines
- HACCP Plans
- Personnel
- Management, Trace back, Sampling and Testing of non-conforming Dairy Product
- Requirements generally based on Codex Programme

# Product Testing and Monitoring

- Operator sampling and testing programmes to confirm conformance are managed under the RMP
- Based on HACCP & stipulated product safety limits
- Market specific requirements are additional
- MPI administered monitoring programmes confirm integrity of the framework:
  - National Chemical Contaminants Programme (NCCP)
  - Independent Verification Programme (IVP)

# Monitoring

## Chemical Residue Monitoring

- Residue monitoring occurs under the National Chemical Contaminants Programme (NCCP) administered by MPI
- NCCP assesses raw milk at the farm and dairy products to:
  - confirm that Good Agricultural Practices are being followed
  - ensure the Regulatory framework is effective
  - ensure market requirements are being met, and
  - identify emerging hazards
- provides an assurance of at least 99% compliance with 95% confidence
- the programme consists of random monitoring, targeted surveillance & periodic surveys



# Monitoring - NCCP

## Compounds monitored include:

- Veterinary medicines & agricultural compounds (including pesticides)
- Contaminants – feeds, farm and process environment
- Adulterants
- Chemical elements & radionuclides
- Compounds of New Zealand or international interest



# Monitoring - IVP

## **Independent Verification Programme:**

- Official sampling of dairy products at packing under the scrutiny of the independent MPI recognised verifier
- Samples submitted to MPI nominated laboratories
- Samples tested for microbiological parameters, standard of identity parameters, or hygiene indicators as determined by MPI
- Strong focus on pathogens, and a bias to sample products for infants and young children
- Typically 300 samples collected each dairy season from premises manufacturing for export

# Verification

# The Contestable Verification Environment

- The APA structure makes provision for non-public sector agencies to be recognised by MPI to be to perform specific tasks under the risk management framework
- Using Recognised Agencies delivers the MPI objective to create a framework allowing industry purchasing of evaluation and verification services in a contestable environment, while maintaining government oversight

# Agencies

## **Recognised Agencies**

Must be accredited to ISO 17020 & recognised by MPI

Evaluate risk management programmes (RMPs)

Verify on-going compliance to RMPs & export requirements

Serve as the front line managers in situation of non-compliance and non-conforming product

Have right of access and certain powers provided to them through the RMP

## **Dairy Laboratories**

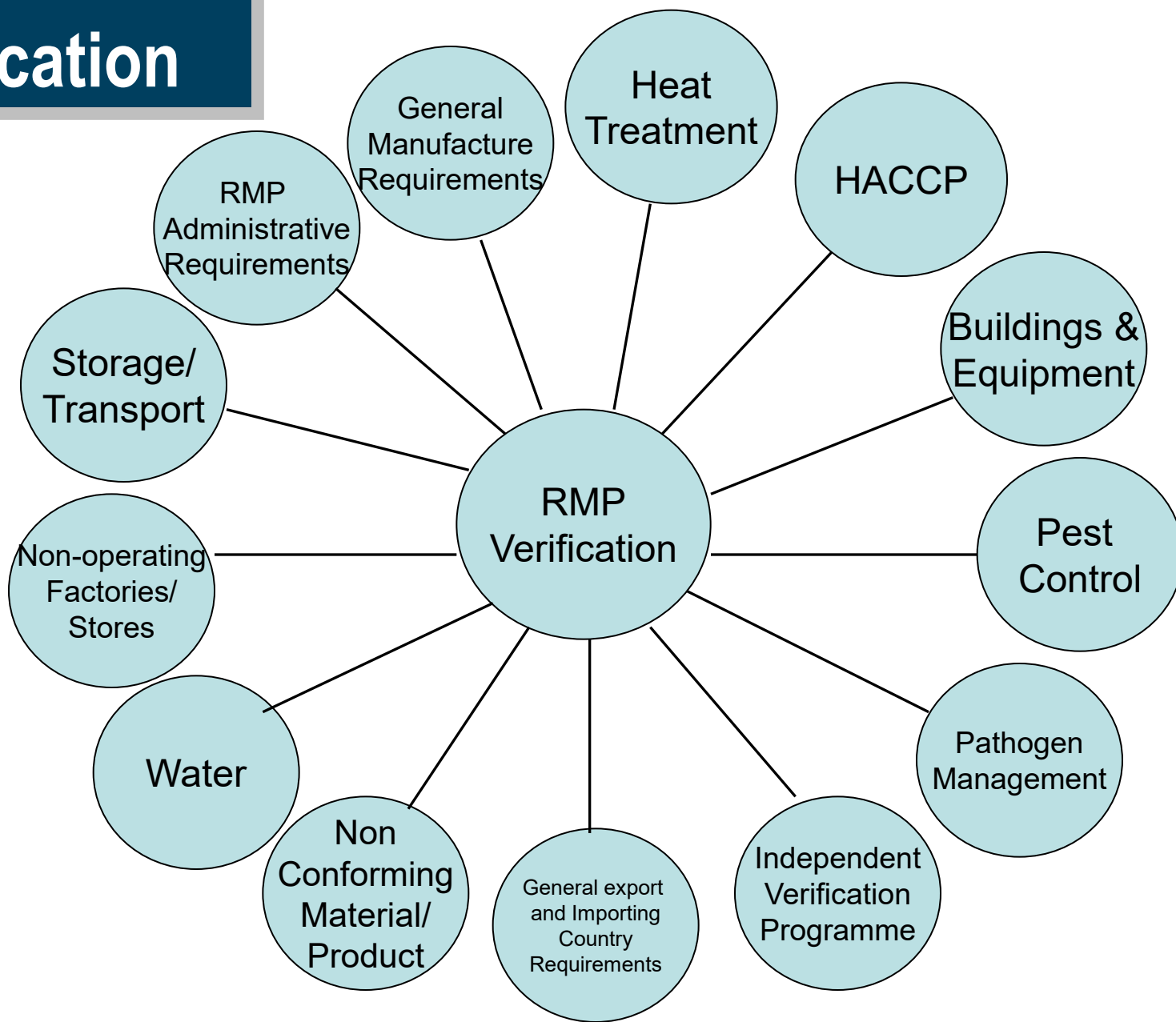
Accredited to ISO 17025 & recognised by MPI

Test Methods must be MPI Approved, though some Markets will specify the methods to be used

# Recognised Agencies

- Third party agencies recognised to act as agents for government
- Verification and other functions eg evaluation of RMPs, verification of export requirements
- Have legal duties
- Cannot apply sanctions
- Must report to MPI – tool used by MPI to monitor industry and agency performance

# RMP Verification



# Non Conformance Management

## Animal Products (Dairy) Regulations 2005

- Section 5 – Non-conforming dairy material or dairy product
- RMP operators must follow the procedures specified by the DG or obtain consent from the DG before disposing of any dairy material or dairy product that is non-conforming.



# MPI Audit and Compliance

- The Systems Audit team within Standards audit elements of the regulatory system to assess compliance as well as assessing the suitability and effectiveness of regulatory measures.
- The MPI Compliance and Response Directorate are responsible for the investigation of suspected breaches under the relevant Acts and to initiate prosecutions where appropriate.
- The Animal Products Act 1999 provides statutory powers to enable Animal Products Officers the power of entry, inspection, sampling, analysis, seizure and forfeiture of products.

# Thank you

