



## Medical Devices Essential Principles Checklist

Manufacturers must demonstrate their medical device complies with the Essential Principles. Manufacturers must generate, collate, assess, and maintain scientific and clinical evidence that shows that their devices comply with the Essential Principles. The evidence must be relevant to the device's intended purpose and must be objective, sufficient, and robust.

- The Essential Principles Checklist is a template that may help you:
  - identify the safety and performance requirements that apply to your device
  - document a rationale for any of the safety and performance requirements that aren't relevant
  - summarise the evidence you hold in support of each of the relevant safety and performance requirements
- Completing the checklist could help you:
  - apply for an Australian conformity assessment certificate
  - comply with regulations as part of a post-market review or investigation
  - to determine if you have considered and addressed all the requirements with supporting evidence

### Instructions

- Complete this checklist for each new medical device conformity assessment application
- Provide an updated checklist if applicable, for change applications relating to your medical device.
- The checklist must be applicable to the subject medical devices or may be completed for a family or group of devices.
- Complete the checklist as outlined in the table instructions below

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### Table instructions

Reference to Therapeutic Goods (Medical Devices) Regulations 2002 Schedule 1, Part number and title		Applicable	Not applicable <i>Justification must be provided below</i>	Medical Devices / IVD standard or other procedure applied	Evidence of compliance <i>Include direct reference to documents within the manufacturer's quality system</i>
EP Reg.#	Title of the Essential Principle				
EP.# Reg.#	The Essential Principle requirement is contained here.	<input type="checkbox"/> Tick this box if the Essential Principle requirement is applicable to the device / family of devices. Complete the two columns on the right.	<input type="checkbox"/> Tick this box if the Essential Principle requirement is not applicable to your device / family of devices. Then provide justification.	List here the standards / procedures applied with justification (include number / title / version). These include (but are not limited to): <ul style="list-style-type: none"> <li>• EN ISO international, local standards</li> <li>• company procedures</li> <li>• TGA Medical Device Standard Order</li> <li>• TGA Conformity Assessment Standard Order</li> <li>• TGA Guidance document</li> </ul>	This column is to contain direct reference to documents within the manufacturer's quality system such as: <ul style="list-style-type: none"> <li>• study results</li> <li>• test reports</li> <li>• design outputs</li> </ul> identified by number / title / version

# Medical Devices Essential Principles Checklist

**J Elliott**



**Medical Devices Essential Principles Checklist:**

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## **Table of Contents Medical Devices Essential Principles Checklist**

1. Understanding the eBook Medical Devices Essential Principles Checklist
  - The Rise of Digital Reading Medical Devices Essential Principles Checklist
  - Advantages of eBooks Over Traditional Books
2. Identifying Medical Devices Essential Principles Checklist
  - Exploring Different Genres
  - Considering Fiction vs. Non-Fiction
  - Determining Your Reading Goals
3. Choosing the Right eBook Platform
  - Popular eBook Platforms
  - Features to Look for in an Medical Devices Essential Principles Checklist
  - User-Friendly Interface
4. Exploring eBook Recommendations from Medical Devices Essential Principles Checklist
  - Personalized Recommendations
  - Medical Devices Essential Principles Checklist User Reviews and Ratings
  - Medical Devices Essential Principles Checklist and Bestseller Lists
5. Accessing Medical Devices Essential Principles Checklist Free and Paid eBooks
  - Medical Devices Essential Principles Checklist Public Domain eBooks
  - Medical Devices Essential Principles Checklist eBook Subscription Services
  - Medical Devices Essential Principles Checklist Budget-Friendly Options

6. Navigating Medical Devices Essential Principles Checklist eBook Formats
  - ePub, PDF, MOBI, and More
  - Medical Devices Essential Principles Checklist Compatibility with Devices
  - Medical Devices Essential Principles Checklist Enhanced eBook Features
7. Enhancing Your Reading Experience
  - Adjustable Fonts and Text Sizes of Medical Devices Essential Principles Checklist
  - Highlighting and Note-Taking Medical Devices Essential Principles Checklist
  - Interactive Elements Medical Devices Essential Principles Checklist
8. Staying Engaged with Medical Devices Essential Principles Checklist
  - Joining Online Reading Communities
  - Participating in Virtual Book Clubs
  - Following Authors and Publishers Medical Devices Essential Principles Checklist
9. Balancing eBooks and Physical Books Medical Devices Essential Principles Checklist
  - Benefits of a Digital Library
  - Creating a Diverse Reading Collection Medical Devices Essential Principles Checklist
10. Overcoming Reading Challenges
  - Dealing with Digital Eye Strain
  - Minimizing Distractions
  - Managing Screen Time
11. Cultivating a Reading Routine Medical Devices Essential Principles Checklist
  - Setting Reading Goals Medical Devices Essential Principles Checklist
  - Carving Out Dedicated Reading Time
12. Sourcing Reliable Information of Medical Devices Essential Principles Checklist
  - Fact-Checking eBook Content of Medical Devices Essential Principles Checklist
  - Distinguishing Credible Sources
13. Promoting Lifelong Learning
  - Utilizing eBooks for Skill Development
  - Exploring Educational eBooks
14. Embracing eBook Trends
  - Integration of Multimedia Elements

- Interactive and Gamified eBooks

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web upon comparison the mrna is found to contain 1 000 fewer bases than the dna sequence did the geneticist isolate the wrong dna a yes mrna is made from a dna template and should be the same length as the gene sequence b yes the mrna should contain more bases than the dna sequence because bases flanking the gene are also transcribed c

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