



INTELLECTUAL PROPERTY
OFFICE OF THE PHILIPPINES

QUALITY MANAGEMENT SYSTEM

Country Report

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27 June 2018

Outline



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- Quality Management System, its Purpose and Benefits
- Patent Quality Review Workflow
- Structure and Role of Quality Management Division
- Documents to be Used for Checking Examination Quality
- Objectives of Patent Quality Manual
- Patent Quality Review Standards
- Sampling Method and Product
- Corrective and Preventive Mechanism

Quality Management Systems



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- IPOPHL:
 - **ISO 9001:2008 Certified** since 2013 which covers the process of granting Patents and registration of Utility Models, Industrial Design and Trademarks
 - **ISO 9001:2015 Certified** since 2017 – transition from ISO 9001:2008

- Bureau of Patents:
 - Bureau of Patents started the development of a **Patent Quality Review System (PQRS)** aimed to assess the quality of issued Office Actions (2013).
 - **Patent Quality Manual (PQM)** was drafted in 2016 and its latest revision was on September 2017.

IPOPHL Quality Policy



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- We strive to foster an environment where IP is created, protected, utilized and enforced.

- We support the creation of a highly-motivated, competent, and cohesive workforce committed to serve with professionalism, transparency, accountability and integrity.

- We are committed to continuously improve our quality management system in order to provide the highest level of satisfaction among our stakeholders.

Bureau of Patents' Quality Commitment



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- We commit to an environment where Patent is protected with fairness, transparency & consistency.

- We provide our staff with knowledge and skills to strengthen competency.

- We dedicate ourselves to continually improve our Patent Quality Examination Standards in order to provide the highest level of satisfaction among our stakeholders.

Purpose of Quality Management System



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- Ensure the quality of search and examination reports.

- Provide reasonable and consistent assessment of search and examination reports.

- Determine the extent of conformity of individuals and groups with the established guidelines.

- Improve the competency of patent examiners.

- Recognize the training needs of patent examiners.

Benefits of Quality Management System



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MANAGEMENT

- Determine what's really going on within the organization, which will allow for more objective decision making.
- Discover where failures occur, enabling the containment of these problems and initiation of corrective actions.
- Identify where resources should be directed.
- Learn which processes and personnel are particularly effective, resulting to recognition.

EXAMINER

- Identify training needs.
- Improvement of technical skills.
- Learn from inaccuracies.
- Increase the efficiency and quality of examination.

3 Types of Quality Check



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• **ISO 9001:2015**

- Covers the process of granting patents and registration of utility model, industrial design and trademarks

• **IN-PROCESS QUALITY CHECK**

- Quality check within the examining division
- 3-Person Team (3-PT)

• **PATENT QUALITY REVIEW SYSTEM**

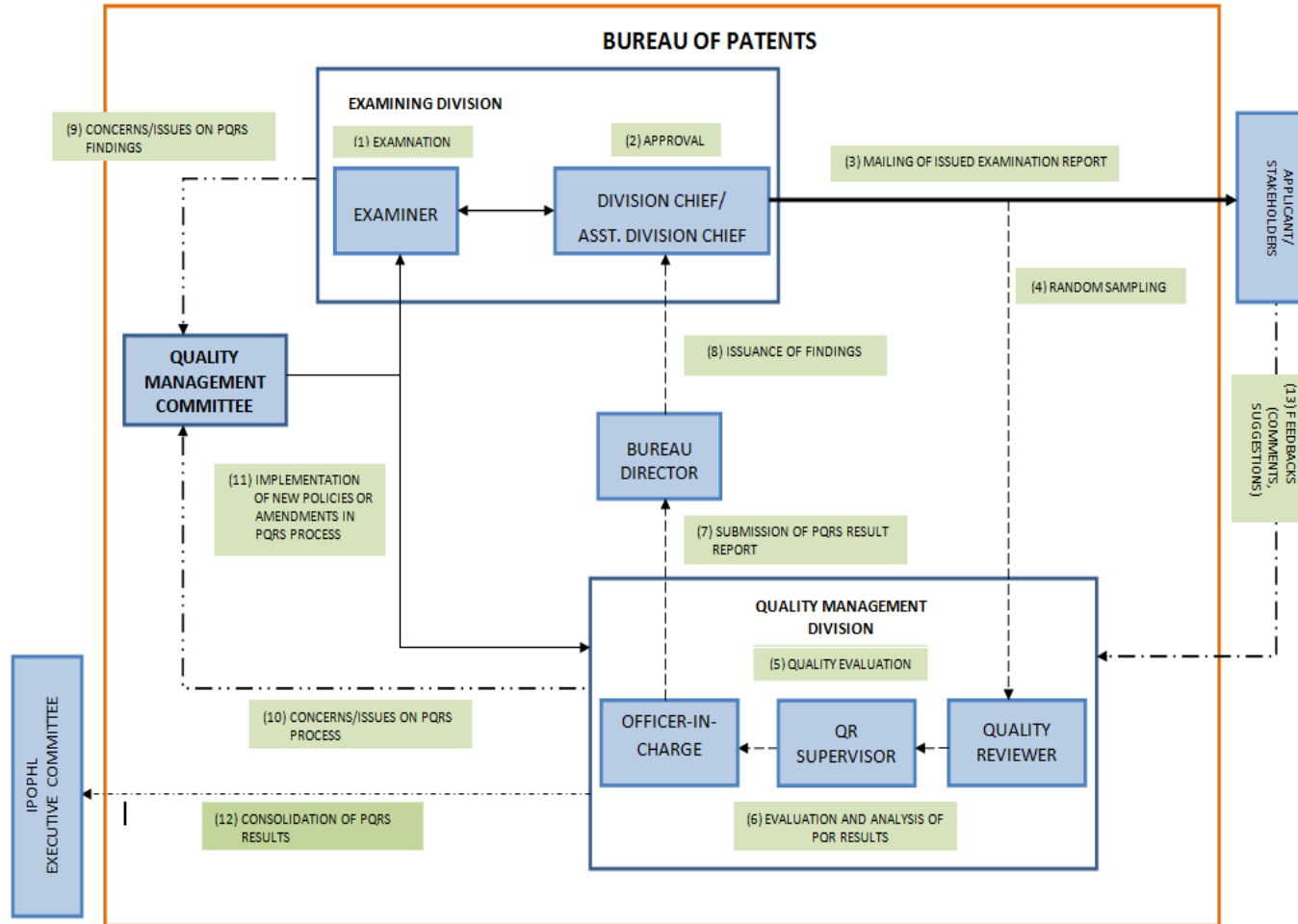
- Random sampling of examination reports and reviewed by the Quality Management Division (QMD)

Patent Quality Review Workflow



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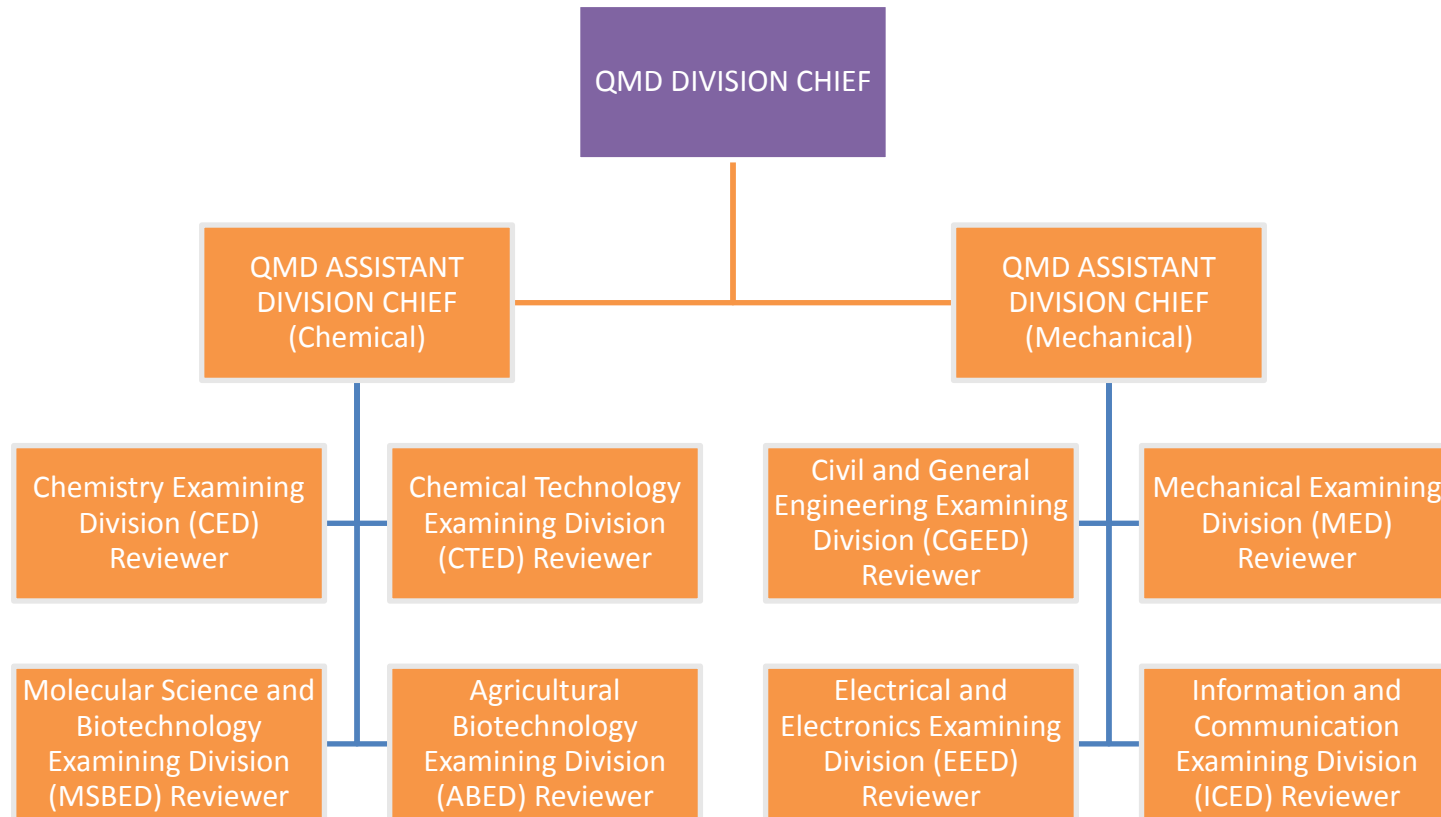
PATENT QUALITY REVIEW WORKFLOW



Structure of Quality Management Division



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Role of Quality Management Division



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- Monitor, maintain and improve the quality of examination and the quality standards.
- Determine the extent of conformity of the examination with the specified standards.
- Determine the effectiveness of the established process.
- Address concerns/issues in examination or process of examination that may occur.

The Quality Reviewer



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- **Qualifications:**

- 5 years of minimum experience in substantive examination and/or has demonstrated high quality of work products for the past 3 years.
- With a performance rating of VS (Very Satisfactory) for at least 2 consecutive years.

- **Target:**

- 10 search and examination reports per month

Role of Quality Reviewers



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- Review the examination report with confidentiality and discretion.
- Evaluates whether the examination report satisfies the quality requirements.
- Evaluates the establishment of reason on patentability.
- Fill out and prepare the patent quality review standards checklist and report form.

Role of Assistant Division Chiefs



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- Evaluate and review the Patent Quality Review Result Form submitted by the Quality Reviewers.

- Serves as the quality check of the Quality Reviewers.

Role of Division Chief



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- Provide a monthly report to the Bureau Director which will highlight the number of conformity and non-conformity findings.

- Identify any particular issue on non-conformity findings that needs immediate attention.

- Evaluate the Examiner's Monthly Rating, Division Rating and the Bureau Rating.

Documents to be Used for Checking Examination Quality



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- Intellectual Property Code (Republic Act 8293)
- Implementing Rules and Regulations (Revised 2017)
- Manual of Patent Examination Practice (Revised 2017)
- Patent Quality Manual (Revised September 2017)
- Others:
 - Guidelines on the Examination of ICT and CII (January 2018)
 - Guidelines on Examination of Biotechnological Applications (January 2018)
 - Guidelines on the Examination of Pharmaceutical Applications involving known Substances (Revised January 2018)

Objectives of Patent Quality Manual



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- To establish patent quality review standards policies, procedures and practices on all the examining and support divisions of Bureau of Patents

- To improve continually the standard, policies, procedures and practices in patent search and examination across all divisions

- To define the role of the Quality Management Division and examining divisions in the patent quality review standards process

Patent Quality Review Standards



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S1 • Patentability

S2 • Searching

S3 • Formality Requirements

S4 • Presentation of Report

*Applicable to Substantive Examination, Search and Written Opinion and Formality Examination

Patent Quality Review Standards



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(S1)

Patentability

- Technical Nature & Exclusions
- Unity of Invention
- Clarity/Support
- Novelty
- Inventive Step
- Industrial Applicability
- Amendments (No New Matter)

(S2)

Searching

- Original Search
- Non-Original Search
- Prior Art

(S3)

Formality Requirements

- Contents of the Application
- Schedule of Fees

(S4)

Presentation of Report

- Completeness of Examiner's Action
- Timeliness
- Documentation

Patent Quality Review Standards – Report Form



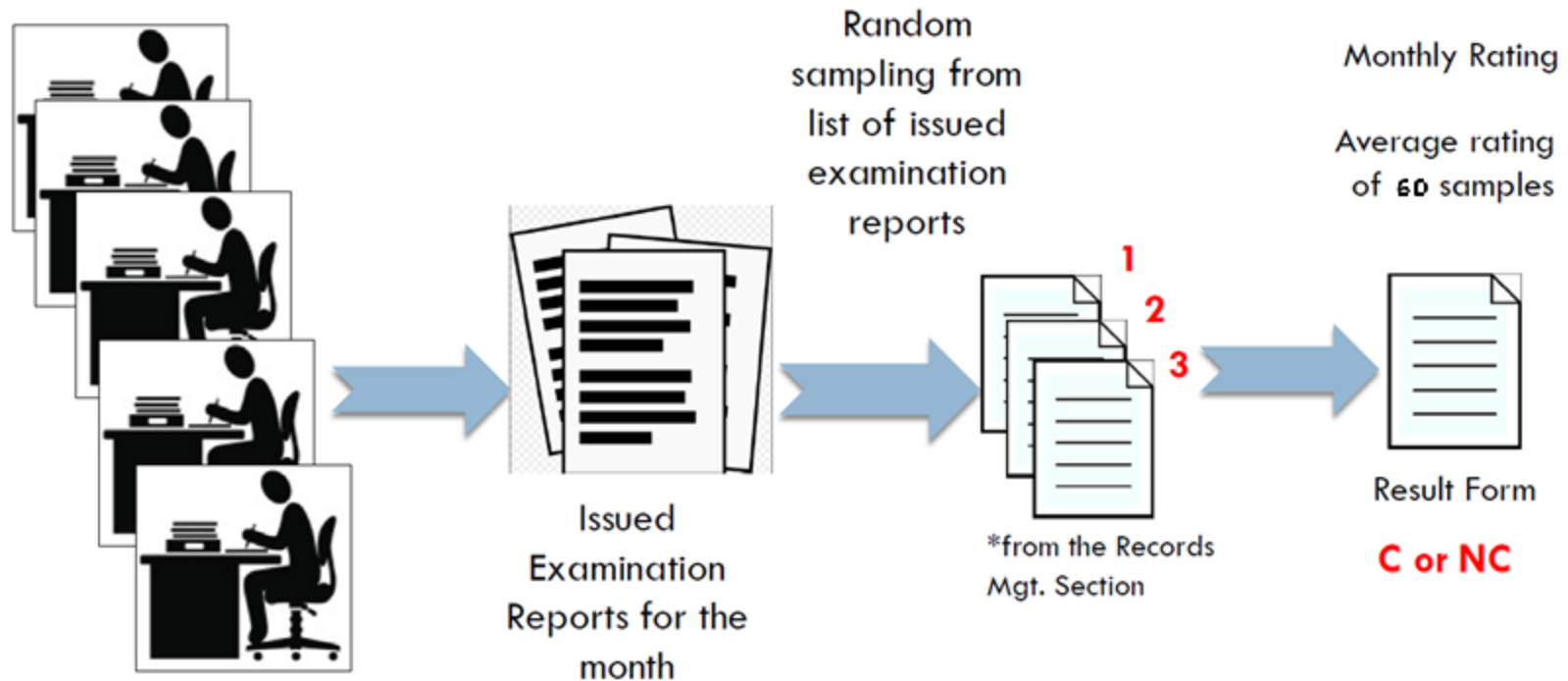
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N.B.: Please return this report to the QMD to acknowledge or respond to the findings within one (1) month. The Examiner's comments and suggestions are hereby solicited for the continuous development and improvement of the Patent Quality Review.	
DIRECTOR	DATE
DIVISION'S RESPONSE	
PREPARED BY:	ACKNOWLEDGED BY:

Sampling Method



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Examination Reports: Substantive Examination, Formality Examination, Search Report

Sampling Method



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- 10 examination reports are reviewed per quality reviewers every month

- Examination reports shall be: search and written opinion; substantive examination report; or formality examination report

- At least 2 examination reports of each patent examiner shall undergo quality review

- Examination reports are randomly selected

Sampling Product



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Patent Quality Review Standards Checklist

Application No. :	I-2010-	Date Issued:	
Examiner Name:		Division:	
Quality Reviewer Name:		Supervisor:	
Month Mailed:	November (2017)	Month Reviewed:	

Type of Examination Report Subsequent Substantive Examination Report

S1	PATENTABILITY	Confor	REMARK
Statutory Basis	Description	C/NC/	
11	Rule 200, 201 & 202	C	
12	Rule 604	C	
13	Rule 415, 405 & 406	C	
14	Rule 203	C	
15	Rule 206	C	
16	Rule 208	C	
17	Rule 916 & 919	NA	
S2	SEARCHING	Confor	REMARK
Statutory Basis	Description	C/NC/	
2.1	Original Search		
2.1.7	Topographical	NA	
2.2	Non-Original Search		
2.2.1	Accepted	NA	
2.3	Prior Art		
2.3.1	Closest Prior Art	C	
S3	FORMALITY REQUIREMENTS	Confor	REMARK
Statutory Basis	Description	C/NC/	
3.1	Rule 400		
3.1.1	Rule 600, 601, 602	C	
3.1.2	Rule 404	C	
3.1.3	Rule 405, 406		
3.1.3.1	Rule 410	C	
3.1.3.2	Rule 407	C	
3.1.4	Rule 413, 414	C	
3.1.5	Rule 415	C	
3.1.6	Rule 411	C	
3.1.7	Rule 418	C	
3.1.8	Rule 418(i)	C	
3.1.9	Rule 305, 306,	NA	
3.1.10	Rule 421	C	
3.1.11	Other(s)	NA	
3.2	Section 13		
3.2.1	Rule 401	C	
3.2.2	Rule 417, 603(e)	C	
3.2.3	Rule 603(e)	NA	
3.2.4	Rule 305, 306	NA	
3.2.5	Rule 401	NA	
3.2.6	Rule 804	C	
3.2.7	Rule 1000	NA	
3.2.8	Rule 1100	C	
3.2.9	Issuance of Letters Patent Certificate	NA	
3.2.10	Other(s)	NA	
S4	PRESENTATION OF REPORT	Confor	REMARK
Statutory Basis	Description	C/NC/	
4.1	Rule 908		
4.1.1	Template	NC	The template should be in the form of a subsequent substantive examination report since the office had already issued two office action (first and subsequent) pertaining to the patentability of the claims.

4.1.2	Correct details and formats	NC	<p>Bibliographic Data: The corresponding IPO box number for applicant's agent was not listed in the 'IPO BOX NO.' field.</p> <p>Acknowledgment: Reassignment of present application to new examiner must be indicated before proceeding to the examination.</p> <p>Basis of the Report: The listed number of pages of specification '1-9' is incorrect. The correct number of pages is '1-17'. Pages should include abstract, description, claims, and drawing sheets.</p> <p>Documents Cited: DI should be listed as obtained by the previous examiner.</p>
4.1.3	Free from	C	
4.1.4	Free from frequent spelling, typographical, or grammatical error	C	
4.1.5	Other(s)	NA	
4.2	Timeliness	C	
4.3	Documentation		
4.3.1	File Wrapper	C	
4.3.2	IPAS Data Entries	C	
OPPORTUNITY FOR IMPROVEMENT (OFI)			
S1: PATENTABILITY			
S2: SEARCHING			
S3: FORMALITY REQUIREMENTS			
S4: EXAMINATION REPORT			

		Quality Reviewer	
		Date	
		QMD OIC / Supervisor	
		Date	

Sampling Product



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Patent Quality Review Standards Report				
Application No. :		1-2010-000		
Examiner Name:		Division:		
Month Mailed:	November (2017)	Supervisor:		
Type of Examination Report:	Subsequent Substantive Examination Report – P1			
Rating:	S1:	C	Opportunity for Improvement	
	S2:	C	S1: <input type="checkbox"/>	S3: <input type="checkbox"/>
	S3:	C	S2: <input type="checkbox"/>	S4: <input type="checkbox"/>
	S4:	NC	Please see comments below	
DESCRIPTION OF FINDINGS				
Below are the observations noted on the above identified Examination Report, stated as follows:				
S1: PATENTABILITY				
The examiner complied with the patent quality review standards for patentability.				
S2: SEARCHING				
The examiner complied with the patent quality review standards for searching.				
S3: FORMALITY REQUIREMENTS				
The examiner complied with the patent quality review standards for formality requirements.				
S4: PRESENTATION OF REPORT				
4.1 Completeness of Examiner's Action				
4.1.1 Template				
The template should be 'IPOP/HL-BOP-INV-FR-02' for final substantive examination report since the office had already issued two office action (first and subsequent) pertaining to the patentability of the claims.				

4.1.2 Correct details and formats	
Bibliographic Data: The corresponding IPO box number for applicant's agent was not listed in the 'IPO BOX NO.' field.	
Acknowledgement: Reassignment of present application to new examiner must be indicated before proceeding to the examination.	
Basis of the Report: The listed number of pages of specification '1-9' is incorrect. The correct number of pages is '1-17'. Pages should include abstract, description, claims, and drawing sheets.	
Documents Cited: D1 should be listed as obtained by the previous examiner.	
N.B.:	
Please return this report to the QMD to acknowledge or respond to the findings within one (1) month.	
The Examiner's comments and suggestions are hereby solicited for the continuous development and improvement of the Patent Quality Review.	
DIRECTOR	DATE

Corrective and Preventive Mechanisms



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• **Corrective Mechanism**

- Issuance of subsequent examination report or re-examination depending on the nature of the non-conformity.

• **Preventive Mechanism**

- Non-conformity issue shall be discussed in the Quality Management Committee (QMC).
- New policies or amendments in the PQRS Process shall be formulated by the QMC for implementation to the Bureau.



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Thank You!