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AMENDMENT RECORDS

No.	Date	Remarks	Revision No.	Approved by
1	03/04/14	Establishment	00	Director, CQA
2	01/04/19	Review the whole contents	01	Director, CQA
3	23/11/21	Review on template of the SOP; item 4.0	02	Director, CQA
		Definitions/ Abbreviation; 7.0 Description;		
		and 8.0 Records		
4	01/07/22	Review on the error of the of the document	03	Director, CQA
		(e.g date format; and abbreviation)		
5	11/10/22	Review on item 5.0 Responsibility; 7.0	04	Director, CQA
		Description		
6	11/06/24	Review on the flow chart and description of	05	Director, CQA
		the working procedure		



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1.0 OBJECTIVE

This SOP describes the process of controlling the documents in UTS and to ensure these documents are being maintained systematically.

2.0 SCOPE

This SOP covers all controlled documents in UTS which are registered under QMS except academic programmes regulated by professional or regulatory bodies.

This SOP applies to all departments and schools that developed and maintained the controlled documents registered under QMS.

3.0 REFERENCES

3.1 ISO 9001: 2015 QMS Manual

3.2 SOP of Control of Record

3.3 SOP of Risk Management

3.4 Other Requirements

4.0 DEFINITIONS / ABBREVIATIONS

UTS : University of Technology Sarawak

CQA : Centre for Quality Assurance
SOP : Standard Operating Procedure
QMS : Quality Management System

HoDs : Head of Departments

DC : Document Controller of CQA

DDC : Department Document Controller

SDC : School Document Controller



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5.0 RESPONSIBILITY

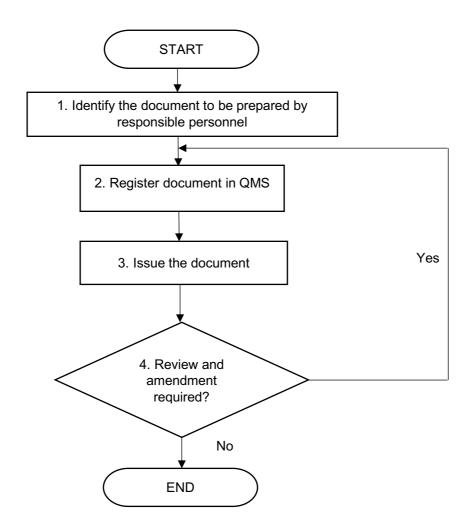
- 5.1 The Director of CQA have the authority to approve the establishment and amendment of controlled document (e.g. SOP, forms, policy, etc)
- 5.2 HoDs and Deans have the authority to verify the establishment and amendment SOP.
- 5.3 HoDs and Deans are responsible to follow and adhere to this SOP.
- 5.4 The director of CQA is responsible to ensure that this SOP is adhered to.
- 5.5 CQA to file and record the 'Master Copy' of the controlled document.
- 5.6 DC to upload the 'Control Copy' of the controlled document in the CQA online platform.

6.0 PROCEDURE

6.1 Refer to the process flow chart.



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7.0 DESCRIPTION

No		Description	Person in Charge	Document
1	Identify the	document to be prepared by	DDC	Controlled
	responsible	personnel.	SDC	Document
	1.1 The Qua	1.1 The Quality Manual and SOP must have the		
	title, doc	ument no., revision no., date, and		Document Change
	page nur	nber.		Form (UTS-CQA-
	1.2 The sup	oporting document related to SOP		P09-DC)
	such a	s the forms must have document		
	no., rev	rision no. and effective date.		
	1.3 Guidelir	ne of numbering the SOP:		
	Doc. No.	University's name: UTS		
		Department's abbreviation: CQA		
		P: Procedure		
		01: Use 2 digits for serializing		
		the SOP starting with "00"		
		e.g. UTS / CQA / P01		
	Revision	Use 2 digits for serializing the		
	No.	revision no. starting with " 00 " for		
		new issue		
	Date	Standardized format for the		
		university is "day/month/year"		
		e.g. 07/01/19		
	Page No.	Current page/Total pages		
		e.g. 1 / 8		
	1.4 Guid	deline of numbering the //guideline:		



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	Doc. No.	University's name: UTS		
		Department's abbreviation: CQA		
		P: Procedure		
		01: Use 2 digits for serializing		
		the SOP starting with "00"		
		Document abbreviation: e.g. DC		
		e.g. UTS – CQA – P09 – DC		
		for Document Change form		
	Revision	Use 2 digits for serializing the		
	No.	revision no. starting with "00" for		
		new issue		
	Effective	Standardized format for the		
	Date	university is "DAY MONTH		
		YEAR"		
		e.g. 07/01/2019		
	docur (UTS- • For no throug depar minut	tment/ school level (e.g. memo, e meeting, etc).		
2			CQA	Controlled Document
	2.1 Relevant department to make sure all		DDC	
	documents which are defined as a		SDC	Master list of
	controlled document must be registered in			Documented
	Master List of Documented Information and			Information (UTS-
	inform CQA. DDC/SDC to submit all SOPs			CQA-P06-MLD)
	and for	ms to CQA.		



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	2.2 Any amendment of the controlled		
	documents, DDC/SDC to update/record		
	the master list of documented information.		
3	Issue the document	DC	Controlled
	3.1 CQA to file and record the 'Master Copy' of	DDC	Document
	the SOP.	SDC	
	3.2 Controlled document shall be uploaded in		
	CQA online platform		
	(https://cqa.uts.edu.my/quality-		
	documents/) by the DC.		
	3.3 All employees are accessible for 'Read-		
	Only'. Any document printed-out from		
	website are considered 'Uncontrolled		
	Copy'.		
4	Review and amendment required?	HoD	Controlled document
	4.1 If yes, process owner to request the	Dean	
	amendment of controlled documents using	Process owner	Supporting
	Document Change Form (UTS-CQA-P09-	DDC	document (e.g.
	DC). Then, repeat the process start from	SDC	memo, minute
	process no. 2 (register document in QMS).	Involved	meeting, etc)
	Any amendment of the controlled	personnel	, ,
	document must gone through meeting		Master list of
	at department/ school level		Documented
	A 'Master Copy' which is obsolete shall		Information (UTS-
	be moved to Archive for reference.		CQA-P06-MLD)
	Other obsolete documents shall be		,
	removed from the distribution list and		Document Change
	prevented from being used.		Form (UTS-CQA-
	provented from being used.		P09-DC)
1			. 33 23,



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,	•	Any amendment of the controlled
		documents, DDC/SDC to update/record
		the master list of documented
		information.
	4.2	If no, end of process.

8.0 RECORDS

No	Title / Records	Location / Responsibility	Retention Period
1	Master list of Documented Information (UTS-CQA-P06-MLD)	Departments Schools CQA	Permanent
2	Obsolete Document Log (UTS-CQA-P06-ODL)	CQA	Permanent
3	Document Change Form (UTS-CQA-P09-DC)	Departments Schools CQA	Minimum of 5 years, unless stated specifically in the procedure
4	Controlled Documents	Departments Schools	Permanent
5	Obsolete Documents	CQA	1 year