

GUIDANCE ON CENTRALIZATION OF BLOOD DONATION TESTING AND PROCESSING



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CONTENTS

Pref	race	v
Ackr	nowledgements	vi
List	of abbreviations and acronyms	viii
Glos	ssary	ix
Exec	cutive summary	xii
Chai	pter 1. Introduction	1
1.1	Purpose and scope of the guidance	
1.2	Target audience	
1.3	Background	
Chap	pter 2. Potential benefits and risks in centralization of blood donation testing and processing	7
2.1	Potential benefits	7
2.2	Potential challenges and risks	10
Chai	pter 3. Key considerations in centralization of blood donation testing and processing	13
3.1	Understanding the need for centralization	
3.2	Structure of the national blood system	
3.3	Which functions should be centralized?	
3.4	Legislative considerations	17
3.5	Good governance	18
3.6	Blood regulation	18
3.7	Organizational considerations	21
3.8	Quality systems	22
3.9	Operational considerations	23
Cha	pter 4. Steps in planning the setting up of a blood establishment to centralize activities	27
4.1	Assessing the current status	
4.2	Blood establishment development plan	28
4.3	Organizational structure	
4.4	Infrastructure	
4.5	Finance and funding	
4.6	Partnerships and agreements	
4.7	Communication	34
4.8	Emergency preparedness.	35

Chap	ter 5. Practical considerations for a blood establishment that will perform centralized blood	
testii	ng and processing	36
5.1	QMS and GMP	36
5.2	Risk management and safety	37
5.3	Organization, roles and responsibilities	37
5.4	Infrastructure	38
5.5	Personnel and training	45
5.6	Funding	46
5.7	Information management systems.	46
5.8	Establishing relationships with stakeholders.	49
5.9	Regulatory aspects	50
5.10	Quality aspects	50
Chap	ter 6. Monitoring and evaluation	52
6.1	Responsibilities for monitoring and evaluation	52
6.2	Monitoring and evaluation	53
6.3	Frequency of evaluation	54
Pofoi	vances	55

PREFACE

The purpose of a national blood system is to ensure timely access to quality, safe and sufficient supplies of blood and blood components for all the patients needing transfusion. However, many low- and middle-income countries are still unable to provide sufficient supplies of whole blood and blood components for patients at all times or in all geographical locations. Vulnerable groups in these countries, such as women with peripartum haemorrhage and children with anaemia from malaria, are often the worst affected by insufficient or unsafe blood supplies. A major barrier to the achievement of universal access to safe blood transfusion in these countries is their ineffective and inefficient blood supply systems. These systems are often fragmented, composed of many small-scale blood establishments, operated by multiple players and have varying levels of performance owing to resource and infrastructure limitations.

Experience in many countries has demonstrated that a nationally coordinated and effectively regulated network of blood establishments, in which key functions are centralized, has many advantages. Centralization optimizes use of resources, reduces overall costs, promotes compliance with quality and safety standards, improves patient access to the most suitable blood components for transfusion, and enhances resilience in emergency situations that affect blood supply or safety. The World Health Organization's *Action framework to advance universal access to safe, effective and quality-assured blood products 2020–2023* recommends countries to consolidate blood donation testing and processing in appropriate facilities. Such facilities should have effectively implemented quality systems to overcome the shortcomings that often exist in decentralized blood systems that are highly fragmented and to achieve the strategic objective of building functioning and efficiently managed blood services.

Guidance on centralization of blood donation testing and processing provides a strategic framework to assist Member States in deciding whether to centralize blood donation testing and processing and how to choose the most suitable centralization model for the unique characteristics of the blood system of each country. The guidance document explains the key steps in planning a blood establishment to centralize activities. It also offers practical guidance on implementing quality, risk management and information management systems, and on managing the infrastructure, human and financial resources necessary for a blood establishment that will perform centralized blood donation testing and processing.

Centralization of blood donation processing can play an important role in increasing the availability of quality plasma for fractionation. This guidance is complementary to the *Guidance on increasing supplies of plasma-derived medicinal products in low- and middle-income countries through fractionation of domestic plasma*. The guidance document provides a strategic framework to assist Member States in increasing their volume of quality plasma suitable for fractionation to help address unmet needs for plasma derived medicinal products.

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