## Achieving excellence Transforming Biomedical Research in clinical trial reporting

A manual for European universities and university hospitals

Workflow

	FIRST STEPS	PAST & PRESENT TRIALS	FUTURE TRIALS
REGISTRY MANAGER	1 Designate manager	1 Old EudraCT trials	[Manage registry data]
	2 Create overview	2 Recent EudraCT trials	
	3 Identify problems	3. All other trials	
TASKFORCE	1 Set up taskforce	1 Strengthen policies 2 Strengthen systems & processes	
	2 Choose model		
	3 Set expectations		

## TABLE OF CONTENTS

INTRODUCTION page 3

Legal, regulatory and funder requirements

Scientific benefits

Registry publication versus journal publication

FIRST STEPS page 4

Designate a registry manager

Create overview of all registry entries

Identify common problems

Set up a taskforce

Choose a model

Set clear expectations Prepare to face two challenges

UPLOAD MISSING CLINICAL TRIAL RESULTS ONTO EUDRACT page 8

Identify trials you are responsible for

Check for data accuracy

Contact your national regulator

Triage trials by their completion date

Contact researchers

Become the 'primary user' of your institution's trials

Upload results as an attachment (older trials only)

Upload results using the full results data set

Overcoming technical problems with uploading results onto EudraCT

page 13

page 14

Useful guidance for uploading results onto EudraCT

## REVIEW YOUR TRIALS LISTED ON OTHER REGISTRIES

Review registry entries

Identify completely unreported trials

Follow up on completely unreported trials

Correct and update registry records

## ADOPTING AND IMPLEMENTING WHO BEST PRACTICES

Why WHO best practices?

Strengthen your policies

Strengthen your systems and processes