

Instructions for printing the ICTRP brochure

The English print version of the brochure can be found at:

www.who.int/ictrp/ictrp_brochure_en.pdf

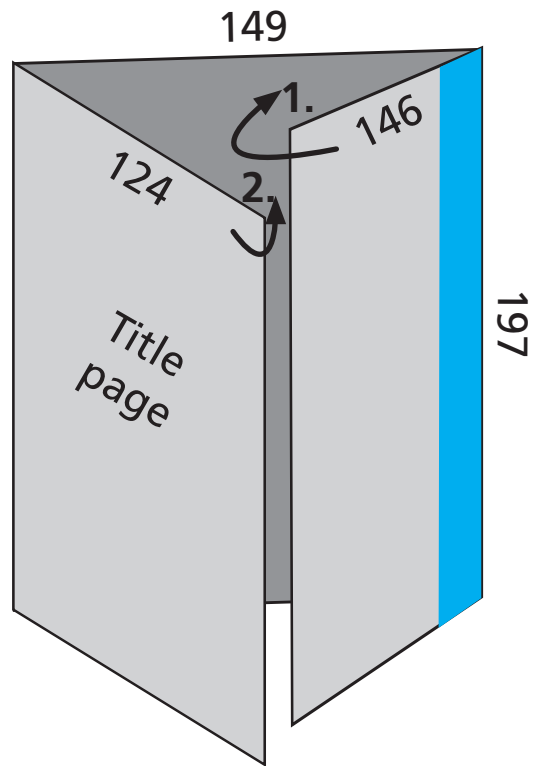
The WHO ICTRP gives permission to print this document under the condition that these guidelines are followed:

- The document should not be altered at all
- No logos should be added to it
- It will not sold or used for commercial purposes
- It should be printed to dimension as laid out on pages 2 and 3 of this document
- It should be cut down neatly to the proper size
- It should be folded as indicated
- It should be printed on thick white mat paper, approximately 200 gsm (grams per square meter)
- You should inform the WHO ICTRP via email (ictrpinfo@who.int) that you are printing it and indicate how it will be distributed

The ICTRP is currently working to translate the brochure into other languages. If you are interested to help us do so, please contact us at ictrpinfo@who.int.

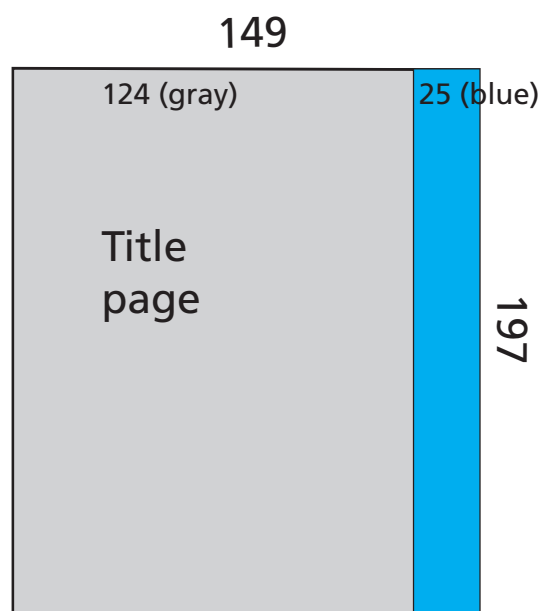
Flyer open: 419 x 197mm

Folded: 2 times (letterfold / parallel fold)



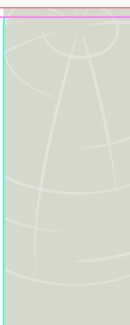



Blue part will be visible when flyer is closed.

Flyer closed: 149 x 197 mm



Fold at
146 mm

Fold at
295 mm

<p>Why we are!</p> <p>The results of clinical trials are not always made available. Worse still, sometimes when a trial is reported only the positive results are made available and not the full picture. In order to counter this lack of transparency countries asked WHO in 2005 to provide a means to link databases that collect information about clinical trials together, provide a single point of access to that information, and make it possible to clearly identify trials and make it easier for patients, families, patient groups and others to access the information.</p> <p>What we are!</p> <p>The ICTRP is a country-led initiative. Clinical trial registration happens when countries seek to improve the transparency of clinical trial research involving nationals of that country, and to be more accountable to the individuals who consent to participate in clinical research, and to better oversee and monitor that research.</p> <p>The ICTRP is a global initiative. It aims to make information about all clinical trials in humans available to all.</p>	<p style="writing-mode: vertical-rl; transform: rotate(180deg);">ICTRP International Clinical Trials Registry Platform</p> <p>What we want to achieve!</p> <ul style="list-style-type: none"> A significant reduction in the gap between what we do and do not know about clinical trials, particularly those conducted in low and middle income countries. The vast majority of currently registered trials are recruiting participants in high-income countries. An increase in the number of countries with either their own national clinical trial registry (meeting WHO standards) or an enforceable policy that clinical trials be registered in a Primary Registry in the WHO Registry Network. An improvement in the quality of registered data. <p>Contact!</p> <p>International Clinical Trials Registry Platform Department of Research Policy and Cooperation World Health Organization Avenue Appia 20n 1211 Geneva Switzerland ictripinfo@who.int www.who.int/ictrip</p>	 <p>International Clinical Trials Registry Platform</p> <p>www.who.int/ictrip</p> 
		

Fold at
124 mm

Fold at
273 mm

<p>What we do!</p> <ul style="list-style-type: none"> Provide an online search portal so that anyone, anywhere can search for information about clinical trials for free. This information is provided by clinical trial registries around the world that meet WHO criteria for content and quality. Support the WHO Registry Network a forum for clinical trial registries around the world to exchange information and work together to establish best practice and ensure the collection of high quality data. Support countries and regions wanting to establish clinical trial registries or policies on trial registration. 	<p>Be Responsible!</p>	<p>What is a clinical trial?</p> <p>A clinical trial is any research study that prospectively assigns human participants or groups of humans to one or more health-related interventions to evaluate the effects on health outcomes. Clinical trials may also be referred to as interventional trials. An intervention can be anything that may have an impact on health, ranging from drugs and surgical procedures through to education, diet and exercise.</p> <p>Clinical trials should be registered in a publicly accessible database before the first participant is recruited. Major scientific journals will no longer publish the findings of a clinical trial unless it was registered before the first participant was recruited in a registry that meets WHO criteria.</p>
<p>The registration of all clinical trials is a scientific, ethical and moral responsibility: www.who.int/ictrip</p>		
<p>The ICTRP aims!</p> <ul style="list-style-type: none"> To improve the comprehensiveness, completeness and accuracy of registered clinical trial data To communicate and raise awareness of the need to register clinical trials To ensure the accessibility of registered data To build capacity for clinical trial registration To encourage the utilization of registered data To ensure the sustainability of the ICTRP 	<p>Be Transparent!</p>	<p>Improved accountability!</p> <p>Our mission is to ensure that a complete view of clinical trial research is accessible to all those involved in health care decision making. Improving transparency will enhance our ability to:</p> <ul style="list-style-type: none"> improve public trust in clinical trial research prevent selective reporting and publication bias meet ethical obligations and protect clinical trial participants Identify research gaps improve recruitment to trials improve trial design, conduct and reporting improve health