

## NHMRC guideline approval process roles and responsibilities

This flowchart provides a summary of the roles of guideline developers, the Office of NHMRC (ONHMRC) and NHMRC Council at each stage of the NHMRC guideline approval process

Stage	Developer	ONHMRC	NHMRC Council
Register (allow 4 weeks for NHMRC to respond)	<ul> <li>Contact NHMRC to discuss eligibility before you start developing your guideline</li> <li>Provide needs analysis and other key details via the Registration Form</li> </ul>	<ul> <li>Assess information provided at registration</li> <li>Write to the NHMRC CEO to seek consideration of the guideline for approval</li> <li>Organise meeting to discuss next steps (if registration is accepted)</li> </ul>	<ul> <li>NHMRC CEO may seek advice from Council before deciding on consideration of the guideline</li> <li>Note the newly registered guideline and its clinical topic area</li> </ul>
2 Develop	<ul> <li>Establish guideline development group</li> <li>Complete systematic reviews of evidence</li> <li>Submit progress report and nominate independent clinical (peer) reviewers 6 months before public consultation</li> <li>Notify NHMRC of public consultation dates</li> </ul>	<ul> <li>Review progress report to identify any major issues or gaps</li> <li>Contact independent clinical (peer) reviewers</li> <li>Contract independent methodological reviewers</li> </ul>	• Note the estimated dates for public consultation and submission for consideration by Council
<b>3</b> Consult (at least 30 days)	<ul> <li>Publish notice inviting public submissions</li> <li>Conduct public consultation (minimum 30 days)</li> <li>Document and respond to submissions</li> <li>Review feedback from independent reviews</li> </ul>	<ul> <li>Commission independent reviews (methodological and clinical expert reviews)</li> <li>Provide de-identified feedback from reviews</li> <li>Help promote public consultation by sending the invitation to Council and publishing a notice on Tracker newsletter</li> </ul>	<ul> <li>Provide feedback on the draft guideline</li> <li>Promote consultation within networks</li> <li>Seek advice from jurisdictions</li> </ul>
<b>4</b> <b>Submit</b> (at least 8 weeks before Council meeting)	<ul> <li>Confirm submission date with NHMRC</li> <li>Submit final draft guideline and other companion documents</li> <li>If asked, join the guideline session at Council meeting as a guest</li> </ul>	<ul> <li>Schedule guideline for NHMRC Council meeting</li> <li>Review and prepare the submission documents for consideration by Council</li> </ul>	<ul> <li>Review and consider the draft recommendations</li> <li>Advise the NHMRC CEO to approve or not approve the recommendations</li> </ul>
5 Publish	<ul> <li><b>Publish</b> guideline within 16 weeks of NHMRC approval</li> <li><b>Make</b> guideline freely available</li> </ul>	<ul> <li>Provide instructions on publishing requirements and use of NHMRC logo</li> <li>Publish approval announcement in NHMRC Tracker newsletter</li> </ul>	• <b>Note</b> that the guideline has been published
<b>6</b> Disseminate	<ul> <li>Implement dissemination plan</li> <li>Engage with stakeholders and promote guideline</li> </ul>	<ul> <li><b>Provide</b> information on NHMRC website</li> <li><b>Support</b> dissemination efforts</li> </ul>	• <b>Support</b> dissemination within Council members' networks where relevant

The content of this resource is based on the *Procedures and Requirements for Meeting NHMRC Standards for Clinical Practice Guidelines*, version 2.0.