

CARE Checklist of information to include when writing a case report



Торіс	Item No	Checklist item description	Reported on Page Number/Line Number	Reported on Section/Paragraph
Title	1	The diagnosis or intervention of primary focus followed by the words "case report"		
Key Words	2	2 to 5 key words that identify diagnoses or interventions in this case report, including "case report"		
Abstract	3a	Introduction: What is unique about this case and what does it add to the scientific literature?		
(no references)	3b	Main symptoms and/or important clinical findings		
	3c	The main diagnoses, therapeutic interventions, and outcomes		
	3d	Conclusion—What is the main "take-away" lesson(s) from this case?		
Introduction	4	One or two paragraphs summarizing why this case is unique (may include references)		
Patient Information	5a	De-identified patient specific information		
	5b	Primary concerns and symptoms of the patient		
	5c	Medical, family, and psycho-social history including relevant genetic information		
	5d	Relevant past interventions with outcomes		
Clinical Findings	6	Describe significant physical examination (PE) and important clinical findings		
Timeline	7	Historical and current information from this episode of care organized as a timeline		
Diagnostic	8a	Diagnostic testing (such as PE, laboratory testing, imaging, surveys).		
Assessment	8b	Diagnostic challenges (such as access to testing, financial, or cultural)		
	8c	Diagnosis (including other diagnoses considered)		
	8d	Prognosis (such as staging in oncology) where applicable		
Therapeutic	9a	Types of therapeutic intervention (such as pharmacologic, surgical, preventive, self-care)		
Intervention	9b	Administration of therapeutic intervention (such as dosage, strength, duration)		
	9c	Changes in therapeutic intervention (with rationale)		

Follow-up and Outcomes	10a	Clinician and patient-assessed outcomes (if available)			
	10b	Important follow-up diagnostic and other test results			
	10c	Intervention adherence and tolerability (How was this assessed?)			
	10d	Adverse and unanticipated events			
Discussion	11a	A scientific discussion of the strengths AND limitations associated with this case report			
	11b	Discussion of the relevant medical literature with references			
	11c	The scientific rationale for any conclusions (including assessment of possible causes)			
	11d	The primary "take-away" lessons of this case report (without references) in a one paragraph conclusion			
Patient Perspective	12	The patient should share their perspective in one to two paragraphs on the treatment(s) they received			
Informed Consent	13	Did the patient give informed consent? Please provide if requested Yes No			

SCARE 2018 Checklist				
Topic	Item	Checklist item description	Page Number	
Title	1	The words "case report" should appear in the title. The title should also describe the area of focus (e.g. presentation,	Page 1, line 1-2	
Key Words	2	3 to 6 key words that identify areas covered in this case	Page 3, line 63	
Abstract	3a	Introduction — Describe what is unique or educational about the case (i.e. what does this work add to the surgical literature, and why is this important?).		
	3b	Presenting complaint and investigations – describe the patient's main concerns and important clinical findings.	Page 3, line 45-	
	3с	The main diagnoses, therapeutics interventions, and outcomes.		
	3d	Conclusion — Describe the main lessons to "take-away" from this case study		
Introducti on	4	Background – summarise what is unique or educational about the case. Give reference to the relevant surgical literature and current standard of care. The background should be referenced, and 1-2 paragraphs in length.	Page 4, line 70- 77	
Patient Informatio n	5a	Demographic details – include de-identified demographic details on patient age, sex, ethnicity, occupation. Where possible, include other useful pertinent information e.g. body mass index and hand dominance.		
	5b	Presentation - describe the patient's presenting complaint (symptoms). Describe the patient's mode of presentation (brought in by ambulance or walked into Emergency room or referred by family physician).	Page 4-5, line 79- 89	
	5c	Past medical and surgical history, and relevant outcomes from interventions		

	5d	Other histories – Describe the patient's pharmacological history including allergies, psychosocial history (Drug, smoking, and if relevant, accommodation, walking aids), family history including relevant genetic information.	
Clinical Findings	6	Describe the relevant physical examination and other significant clinical findings. Include clinical photographs where relevant and where consent has been given.	Page 4-5, line 79-
Timeline	7	Inclusion of data which allows readers to establish the sequence and order of events in the patient's history and presentation (using a table or figure if this helps). Delay from presentation to intervention should be reported.	Page 4-5, line 79- 89
Diagnosti c Assessme	8a	Diagnostic methods – describe all investigations taken to arrive at methods: physical exam, laboratory testing, radiological imaging, histopathology.	
nt	8b	Diagnostic challenges – describe what was challenging about the diagnoses, where applicable, for example access, financial, cultural.	Page 5, line 89-
	8c	Diagnostic reasoning – Describe the differential diagnoses and why they were considered.	100
	8d	Prognostic characteristics when applicable (e.g. tumour staging or for certain genetic conditions). Include relevant radiological or histopathological images in this section.	
Therapeut ic Interventi on	9a	Pre-intervention considerations – if there were patient-specific optimisation measures taken prior to surgery or other intervention these should be included e.g. treating hypothermia/hypovolaemia/hypotension in a burns patient, Intensive care unit treatment for sepsis, dealing with anticoagulation/other medications, etc.	
	9b	Interventions – describe the type(s) of intervention(s) deployed (pharmacologic, surgical, physiotherapy, psychological, preventive). Describe the reasoning behind this treatment	Page 5- 6, line 100- 113

	9c	offered. Describe any concurrent treatments (antibiotics, analgesia, anti-emetics, nil by mouth, Venous thromboembolism prophylaxis, etc). Medical devices should have manufacturer and model specifically mentioned. Intervention details – describe what was done and how. For surgery include details on; anaesthesia, patient position, use of tourniquet and other relevant equipment, prep used, sutures, devices, surgical stage (1 or 2 stage, etc). For pharmacological therapies include information on the formulation, dosage, strength, route, duration, etc. Include intra-operative photographs and/or video or relevant histopathology in this section. Degree of novelty for a surgical technique/device should be mentioned e.g. "first in human". Who performed the procedure - operator experience (position on the learning curve for the technique if established,	Page 5- 6, line 100- 113
	Ju	specialisation and prior relevant training). For example, "junior resident with 3 years of specialised training"	
	9e	Changes – if there were any changes in the interventions, describe these details with the rationale.	
Follow-up and Outcomes	10a	Follow-up – describe 1) When the patients was followed up. 2) Where. 3) How (imaging, tests, scans, clinical examination, phone call), and 4) whether there were any specific post-operative instructions. Future surveillance requirements - e.g. imaging surveillance of endovascular aneurysm repair or clinical exam/ultrasound of regional lymph nodes for skin cancer.	
	10b	Outcomes - Clinician assessed and (when appropriate) patient- reported outcomes (e.g. questionnaire details). Relevant photographs/radiological images should be provided e.g. 12 month follow-up.	Page 6, line 113- 114
	10c	Intervention adherence/compliance - where relevant how well patient adhered to and tolerated their treatment. For example,	

		post-operative advice (heavy lifting for abdominal surgery) or tolerance of chemotherapy and pharmacological agents	
	10d	Complications and adverse events – all complications and adverse or unanticipated events should be described in detail and ideally categorised in accordance with the Clavien-Dindo Classification. How they were prevented, diagnosed and managed. Blood loss, operative time, wound complications, reexploration/revision surgery, 30-day post-op and long-term morbidity/mortality may need to be specified. If there were no complications or adverse outcomes this should also be included.	Page 6, line 113- 114
Discussio	11a	Strengths – describes the strengths of this case	
n	11b	Weaknesses and limitations in your approach to this case. For new techniques or implants - contraindications and alternatives, potential risks and possible complications if applied to a larger population. If relevant, has the case been reported to the relevant national agency or pharmaceutical company (e.g. an adverse reaction to a device)	Page 6-9, line
	11c	Discussion of the relevant literature, implications for clinical practice guidelines and any relevant hypothesis generation.	120- 192
	11d	The rationale for your conclusions.	
	11e	The primary "take-away" lessons from this case report.	
Patient Perspectiv e	12	When appropriate the patient should share their perspective on the treatments they received.	Page5, Line10 3-105
Informed Consent	13	Did the patient give informed consent for publication? Please provide if requested by the journal/editor. If not given by the patient, explain why e.g. death of patient and consent provided by next of kin or if patient/family untraceable then document efforts to trace them and who within the hospital is acting as a guarantor of the case report.	Page 6, line 117 Page 5, line 104

Additional			
Informatio	14	Conflicts of Interest, sources of funding, institutional review	Page 10, line 205-
n		board or ethical committee approval where required.	212

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*As the checklist was provided upon initial submission, the page number/line number reported may be changed due to copyediting and may not be referable in the published version. In this case, the section/paragraph may be used as an alternative reference.