Use of carbon dioxide as an intravascular contrast agent: A review of current literature

Fahad Ali, Muhammad Asif Mangi, Hiba Rehman, Edo Kaluski

Fahad Ali, Division of Cardiology, Department of Medicine and Lehigh Valley Hospital, Allentown, PA 18105, United States

Muhammad Asif Mangi, Hiba Rehman, Department of Pulmonary and Critical Care Medicine, Mayo Clinic, Jacksonville, FL 32224, United States

Edo Kaluski, Division of Cardiology, Department of Medicine, Guthrie Robert Packer Hospital and Guthrie Health Services, Sayre, PA 18840, United States

Author contributions: All authors contributed to the manuscript equally; all authors read and approved the final manuscript.

Conflict-of-interest statement: None of the authors have any conflicts of interest relative to this publication.

Open-Access: This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/

Manuscript source: Unsolicited manuscript

Correspondence to: Fahad Ali, MD, Division of Cardiology, Department of Medicine and Lehigh Valley Hospital, P.O. Box 689, Allentown, PA 18105, United States. drfahadalis@yahoo.com
Telephone: +1-570-8673025

Received: January 11, 2016
Peer-review started: January 14, 2016
First decision: March 7, 2016
Revised: October 9, 2016
Accepted: October 22, 2016
Article in press: October 24, 2016
Published online: September 26, 2017

Abstract

Use of X-ray contrast allows us to differentiate between two or more adjacent structures on radiographic studies. The X-ray contrast agent can be the one with increase X-ray absorption, like iodine and a barium X-ray contrast agent or the one with decrease X-ray absorption like air and carbon dioxide contrast agent. Each contrast agent possesses different risks and benefits in various ways. Carbon dioxide as an intravascular contrast agent can be used as an alternative intravascular contrast agent and has superior results in some cases. In patients with renal dysfunction or iodinated contrast allergy, the use of iodinated Contrast Agent poses the risk of considerable morbidity. Similarly, use of Gadolinium is discouraged in subject with severe renal dysfunction. Use of carbon dioxide (CO₂) as an intravascular contrast, offers an alternative in such patients for certain procedures, as it is not nephrotoxic and it does not incite allergic reactions. It is inexpensive, readily available and due to its unique physical properties, it can be used to image a wide variety of vascular beds and chambers. The aim of this paper is to systemically review the current literature to describe the indications, contraindications, adverse effects, instruments, precautions, latest methodologies and data supporting for the use of CO₂ as a contrast agent.

Key words: Iodinated; Carbon dioxide; Contrast; Vascular; Gadolinium

© The Author(s) 2017. Published by Baishideng Publishing Group Inc. All rights reserved.

Core tip: In patients with renal dysfunction or iodinated contrast allergy, use of iodinated contrast agent poses the risk of considerable morbidity. Similarly, use of gadolinium is discouraged in subject with severe renal dysfunction. Use of carbon dioxide (CO₂) as an intravascular contrast offers an alternative in such patients for certain procedures, as it is not nephrotoxic and it does not incite
allergic reactions. It is inexpensive, readily available and
due to its unique physical properties it can be used to
image a wide variety of vascular beds and chambers. This
article describes the indications, contraindications, adverse
effects, instruments, precautions, latest methodologies
and data supporting for the use of CO₂ as a contrast
agent.

Ali F, Mangi MA, Rehman H, Kaluski E. Use of carbon dioxide
as an intravascular contrast agent: A review of current literature.
World J Cardiol 2017; 9(9): 715-722 Available from: URL:
http://dx.doi.org/10.4330/wjc.v9.i9.715

INTRODUCTION
In medical parlance, contrast is a mean which allows
us to differentiate between two or more adjacent
elements on a radiographic study. There are essentially
two prototypes of X-ray contrast agents: (1) Positive
agents (which increase X-ray absorption: Iodine or
barium based); and (2) negative agents [decrease
X-ray absorption: Air, carbon dioxide (CO₂)]1. In ani-
mal experiments (1940s) and later in human studies
(1950s), CO₂ enabled investigators to delineate both
right and left heart structures. With the introduction
of digital subtraction angiography (DSA) in 1980s, the
image quality improved significantly2. Conditions where
use of iodinated contrast agent (ICA) are precluded
such as impaired kidney functions, dye allergy, CO₂ may
be used as an alternate contrast agent with comparable
results and in some cases superior results3-4.

Physical properties
Understanding of physical properties of CO₂ is central
for its use. Administration of CO₂ needs extreme care.
It is a colorless, odorless and significantly compressible
gas. CO₂ has low molecular weight as compared to
ICA, is less viscous then blood and ICA and due to
this property it can be used to image small collateral
vessels. It displaces the blood in the vessels and acts
as a negative contrast agent. This property creates a
significant gradient between the radiographic density of
the vessel wall and the lumen. DSA technique uses this
difference in the densities to provide a contrast image.
CO₂ is more soluble than oxygen (O₂) and dissolves in
the blood within 2-3 min after injection. When mixed
with water it creates carbonic acid (H₂CO₃) which dis-
sociates into bicarbonate (HCO₃⁻) and hydrogen (H⁺)
ions carried by blood flow to the lungs. Reverse reaction
happens in the lungs where the breakdown product of
H₂CO₃, CO₂ is then exhaled. These chemical reactions
are facilitated by enzymes called carbonic anhydrases.

During its use, monitoring of vital signs is required. Capnography if available would be useful in monitoring
the ventilation.

Administration
There are 3 commonly used methods of administering
CO₂. Preferred method is via automated injectors
(automated CO₂ mmmanders). Hand held syringes have
been used in the past but are not commonly used
now due to increased risk of complications such as air
contamination and explosive over dosage5-6.

Automated CO₂ mmmanders: Have the utility of being
handy, portable, safe and easy to use but their high cost
(approximately 3000 USD)7-9 make them an unpopular
choice.

The modified plastic bag system with O-ring: Is a
preferred method by some experts. Kit packs consisting
of bag, tube and valves are available commercially
(custom waste management kit by Merit Medical,
South Jordan, UT; or Anglo-Dynamics Queensbury,
NY)8,9. The usual source of CO₂ is an Aluminum or steel
cylinder of medical grade CO₂ which is about 99.99% pure,
packed in a series circuit with a valve, a gauge, a
regulator, a diaphragm and an antibacterial filter. A
1500 mL plastic bag with a single port connected with
a low pressure tube and a 2-way stopcock at the distal
end of the tube is then connected to the CO₂ cylinder.
It is then filled and manually purged at least 3 times.
The filled bag is then connected at its 2-way stopcock
end with an O-ring connector which on the other end
is connected with the delivery syringe (20-60 mL). There
is a 1-way valve between the O-ring and the syringe.
The syringe is then connected with another 1-way valve
and then with a 100 cm connecting tube. The distal end
of this tube has one more 1-way valve which is then
connected with a 3-way valve. This 3-way valve can
then be connected with the angiographic catheter. On
the other port an additional syringe for back-bleeding
or eliminating the air from the system can be attached
(Figure 1). To fill the delivery syringe the plunger is
simply retracted. The 1-way valves will allow the CO₂ in
the plastic bag to move into the syringe. The plunger
can then be advanced at the desired rate and amount.
The 1-way valves will allow the gas to move towards
the 3-way valve which can then be adjusted depending
on the ports required to be used. The angiographic
catheter is at times filled with blood which can be
cleared by using the additional syringe attached at the
3-way valve. Forceful boluses of 3-5 mL CO₂ can be
used to clear the catheter from any remaining fluid. The
catheter can then be flushed with 1-3 mL of CO₂ every
2-3 min. All the connections of the circuit need to be
air tight to avoid any air aspiration or embolism. The
plastic bag should be filled enough to remain flaccid as
tightly filled bag may pose risk of overdose due to gas
compression9.

Underwater seal: This is a relatively simple, inex-
pensive and easier method but there may be a slight
risk of air contamination and or inadvertent explosive
administration of CO₂ into the patient[10]. In this system, the CO₂ source is connected to a regulator, a particle filter and a 3-way tap by connecting tubes. One end of 3-way valve has a sliding 2-way valve connected to a 60 cc syringe. The other end of 3-way valve has a tube serving as a simple under water seal by having the other end of the tube dipped in a bowl of saline. When the CO₂ source is turned on and the 3-way valve is on to the syringe, the syringe will get filled without pulling the plunger by the positive pressure of the CO₂ coming from the source. Once the syringe is filled the 2-way valve is turned off and 3-way valve is turned to the under-water seal. Bubbles of CO₂ would be seen in the bowl of saline coming out of the tube’s end. The CO₂ source is then turned off and the 3-way valve is then turned on to the syringe and the water seal. The CO₂ can then be purged through the water seal. This process of filling and purging can be repeated at least 3 times to make sure that there is only CO₂ and no air in the tubing and syringe system. Then the filled syringe along with a 2-way valve turned off, can then be disconnected and attached to the angiography catheter. Right before it is connected to the angiography catheter, the 2-way valve is turned on to release the positive pressure in the syringe to come down to atmospheric pressure. This will avoid explosive administration and or over dosing of pressurized CO₂ in the syringe but at the same time this may create a very small risk of air contamination. Only fully filled syringes should be used while using this method as half-filled syringes when opened to atmospheric pressure will certainly lead to higher risk of air contamination. The innovators of this system also described their experience of 5 years in over 250 patients and no directly related complications were noticed[10].

**Dosage**

Typically 30-40 mL of CO₂ is injected for abdominal aortography or IVC visualization. Twenty to thirty milliliters is used for lower extremity vessels and other aortic branches like celiac, superior mesenteric or renal arteries. The left renal artery which is more posteriorly located can be filled even with 10 mL if injected with patient lying on the right side. Injections can be repeated at approximately 3 min intervals. Thirty to fifty milliliters may be used for runoff studies by injecting at low rate of 10 mL/s.

**Potential uses of CO₂-based angiography (Figure 2)**

The diagnostic accuracy is acceptable in comparison to contemporary ICA and in some conditions such as TIPS, CO₂ is even rendered superior to the ICA.

**Aortic aneurysm repairs**: CO₂ has been used in endovascular repairs of aortic aneurysms[11-14]. A recent prospective study of 72 patients with abdominal aortic aneurysm (AAA) endovascular repair demonstrated that CO₂ has overall sensitivity of 84% and specificity of 72% as compared to ICA as the standard criterion for detection of endoleaks and in patients who are at risk of nephrotoxicity from ICA, CO₂ can be used as an acceptable alternative to ICA[15]. Another study describes the outcomes of CO₂-guided procedures are similar to those which are ICA-guided[16]. Additional benefit of CO₂ use in endovascular repair of AAA is that an accessory catheter which is otherwise required for ICA may not be required for CO₂ injection as it can be administered through the endograft sheath or femoral access sheath[13].

**Aortography**: CO₂ may be used for aortography and for runoff studies in most patients[16]. If needed supplemental ICA imaging may be used in order to obtain additional information. To get the retrograde aortogram, CO₂ may be injected retrograde through the femoral artery by percutaneous catheterization with a 4-Fr end-hold catheter (Cobra-shaped or shepherd hook catheter) or catheters with side-holes (Omni-flush, pigtail, Racquet, multipurpose). Contra-lateral superficial femoral arterial views can also be taken through the same port by moving the catheter into the contralateral superficial femoral artery. For antegrade views micro-catheters of 3-Fr may be used for popliteal, tibial and peroneal arteries. Use of intra-arterial nitroglycerine and or leg elevation may be done for better visualization of smaller vessels such as tibial and plantar branches.

**Renal artery angiography (Figure 3)**: CO₂ can be used in the assessment of renal artery stenosis, aneurysms, AV (arterio-venous) malformations, AV fistulas, renal artery stenting, invading tumors in renal
Uses of intra-vascular carbon dioxide as a contrast agent

Venous
- Inferior vena cava procedures
- TIPS
- Upper extremity venous procedures
- Inferior vena cava imaging

Arterial
- Arterial run-off
- AAA
- Renal arterial procedures
- Mesenteric arterial procedures
- Peripheral arterial procedures
- Arterio-venous fistulography

Arterio-venous fistulography

Neoplasia related procedures
- Chemoembolization of hepatocellular carcinoma
- Chemoembolization of renal cell carcinoma

Figure 2  Potential uses of carbon dioxide angiography. AAA: Abdominal aortic aneurysm.

veins or arteries, renal cell carcinomas, evaluation of transplanted kidney vascular stenosis and for its angioplasty and/or stenting, anastomotic stenosis, diffuse arterial disease related to chronic rejection and AV fistulas after renal transplant biopsy (in which case it may be superior to ICA)\textsuperscript{[17,18]}\textsuperscript{17,18}. In such cases CO\textsubscript{2} may be used as initial imaging modality to get an overview and then small dose ICA may be used for confirmation of the findings\textsuperscript{[11]}\textsuperscript{11}. CO\textsubscript{2} does not adequately fill the distal portion of renal artery very well in a supine patient, as it is located posterior to the aorta. In this situation, the patient may be turned on the side to bring the renal arteries superior with respect to the aorta. Recent studies have also demonstrated the use of CO\textsubscript{2} in combination with intravascular ultrasound for successful vascular stenting. In a study of 18 patients, 27 successful renal artery stenting procedures were done using CO\textsubscript{2} and intravascular ultrasound with good outcomes\textsuperscript{[19,20]}\textsuperscript{19,20}.

**Inferior vena cava imaging:** CO\textsubscript{2} can be used for the placement of inferior vena cava (IVC) filters, IVC venous anomalies and thrombus visualization, recanalization of occlusion and estimation of IVC diameters (accuracy of about 97%). In a study of 50 patients, CO\textsubscript{2} was used for IVC filter placement at the bedside in ICU setting. Only 2 of these patients required additional ICA for better visualization. The study concluded with positive results and favored the use of CO\textsubscript{2} as first line contrast agent in ICU patients requiring IVC filter\textsuperscript{[21]}\textsuperscript{21}.

**Portal vein imaging (portography):** A very important utility of CO\textsubscript{2} is in the delineation of the portal vein anatomy (wedged hepatic venography) during TIPS procedure (Figure 4). CO\textsubscript{2} is found to be superior to ICA for this use and can be used as first line contrast agent for portography\textsuperscript{[22]}\textsuperscript{22}. The reason is buoyancy and low viscosity of CO\textsubscript{2} making it travelling through the sinusoids easily and deeply. In liver transplants, anastomosis can also be visualized using CO\textsubscript{2}. In a study of 16 patients, the utility of CO\textsubscript{2} was compared with ICA for balloon-occluded retrograde trans-venous venography (BRTV) and obliteration (BRTO) for gastric varices and it was found that varices were visualized...
for the following oncological embolization procedures: Embolization of renal cell carcinoma and its metastatic lesions in the bone, hepatocellular carcinoma[28,29], radiofrequency ablation and transcatheter arterial chemoembolization of hepatocellular carcinoma (by using intra-arterial CO\textsubscript{2} for enhancement for ultrasonography guidance)[30], uterine artery embolization in uterine leiomyoma[31]. These procedures can be optimized by using super-selective angiographic techniques with help of micro-catheters of 3 Fr.

**Upper extremity venography:** Can be performed using the CO\textsubscript{2}[32]. It can be useful for AV-fistula formation[33], insertions of trans-venous pacer wires[34], central venous catheters and for the delineation of any atypical vascular anatomy. The preferable site of injection is antecubital vein and a 21 gauge catheter may be used. In a series of 146 AV fistulography procedures using CO\textsubscript{2} as the first line contrast agent, 141 cases required AV fistula intervention and in 115 of these cases intervention was performed successfully using CO\textsubscript{2} alone. Rest of the cases required ICA for various reasons in addition to CO\textsubscript{2} for intervention[35].

For AV fistula assessment, one needs to be careful of not letting CO\textsubscript{2} reflux into arterial system due to potential risk of neurologic sequelae including infarction. Also there is a potential of overestimation of fistula stenosis.

**Gastrointestinal bleeding:** Due to increased compressibility and low viscosity it may be useful in detecting the site of occult bleeding or ongoing blood loss such as the gastrointestinal tract, with higher sensitivity than ICA. CO\textsubscript{2} can also be used in selected angiographies for chronic mesenteric ischemia[36].

**Contrast ultrasonography:** CO\textsubscript{2} can be used to enhance sonography by employing CO\textsubscript{2} microbubbles. In a study where conventional sonography was compared with CO\textsubscript{2} micro-bubble enhanced sonography; the former detected only 6 tumors however with CO\textsubscript{2}-microbubble enhanced sonography 14 tumors were detected and then treated successfully with radiofrequency ablation.
using CO\textsubscript{2}-microbubbles enhanced sonography\cite{37}.

**LIMITATIONS**

Overall CO\textsubscript{2} is a relatively safe agent\cite{38}. In a study of 800 subjects, only one complication of transient colonic ischemia was reported. In another study of 1200 subjects only 7 subjects developed some kind complication. Livedo reticularis, bowel ischemia and renal dysfunction have been described after in 1 patient with CREST syndrome\cite{38} (Table 1).

The adverse effects are primarily either dose related or buoyancy related. Majority of the adverse effects are due to “vapor-lock phenomenon” which result when large amounts of CO\textsubscript{2} are injected or a small amount is injected too frequently with very short intervals causing trapping of CO\textsubscript{2} gas column in the vessel and consequently obstructing the vessel. This may lead to ischemia of the tissues. Cases with transient mesenteric ischemia and ischemic colitis secondary to “vapor lock phenomenon” have been described in the literature. Similar mechanism may potentially precipitate right sided heart failure. Sometimes CO\textsubscript{2} bubbles may accumulate in an aortic aneurysm and may cause blood flow obstruction, leading to tissue ischemia. Even a transient occlusion of inferior mesenteric artery may result in mesenteric ischemia. Typically, this happens with the use of excessive dose of CO\textsubscript{2}. Similarly a vapor lock may happen in the pulmonary artery and this may lead to significant hypotension. Air contamination may also cause vapor-lock phenomenon that is typically worse and more persistent. Usually CO\textsubscript{2} bubbles in the pulmonary artery dissolve within 30 s. If they persist beyond 30 s then either air contamination or CO\textsubscript{2} over-dosage should be suspected and the tubing system should be checked for any air leak. For hypotension secondary to vapor lock phenomenon, patient should be placed in Trendelenburg position or lateral decubitus positions. Aspirating the air using a catheter from the pulmonary artery should also be considered. Due to its buoyancy the visualization of a dependent (inferior or caudal positioned) vessels may be suboptimal (such as visualization of renal arteries in supine position). This problem can be circumvented by putting the patient in lateral decubitus position. CO\textsubscript{2} may get trapped in organs which are non-dependant and cause decreased blood flow or ischemia such as in transplanted kidneys or mesenteric vessels. If we place the patient in lateral decubitus position, the CO\textsubscript{2} may remain trapped in right atrium instead of moving into pulmonary arteries. Changing the body position may help clearance in these situations. Similarly using low volumes of CO\textsubscript{2} with adequate time intervals may help avoiding these adverse effects. Although 100 mL is the recommended maximum volume for arterial use and 50 mL for venous use, by using above mentioned precautions, larger total volume may be used. Vessels which are more anterior such as superior mesenteric artery (SMA), CO\textsubscript{2} is useful in their evaluation, particularly for proximal mesenteric stenosis. For more distal assessment ICA probably provides superior imaging.

Due to its dissolution in blood soon after injection vessels with slower flow may not have adequate visualization and this may lead to overestimation of stenosis. Similarly due to the expansive nature of CO\textsubscript{2} and elasticity of vessels, CO\textsubscript{2} may lead to overestimation of the vessel diameter. This may cause errors in estimation of balloon or stent size during intervention procedures.

The use of CO\textsubscript{2} for cerebral, spinal and or cardiac procedures should be avoided as there is a potential risk of ischemia to vital organs\cite{39}. In animal models neurotoxicity has been reported after cerebral use. For the same reasons, before the use of CO\textsubscript{2} presence of atrial or ventricular septal defects or pulmonary arteriovenous malformation should be ruled out to avoid the risk of paradoxical embolism to CNS and or coronary embolization. CO\textsubscript{2} may also aggravate or worsen pulmonary arterial pressure therefore the use of this agent should be avoided in pulmonary hypertension. There are some relative contraindications for the use of CO\textsubscript{2} for upper extremity which are similar for other uses.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Carbon dioxide</th>
<th>Iodinated contrast agents</th>
</tr>
</thead>
<tbody>
<tr>
<td>Overall sensitivity</td>
<td>Less</td>
<td>Higher</td>
</tr>
<tr>
<td>Overall specificity</td>
<td>Less</td>
<td>Higher</td>
</tr>
<tr>
<td>Nephrotoxicity</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>Allergic toxicity</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>Cost</td>
<td>Low</td>
<td>High</td>
</tr>
<tr>
<td>Ease of administration</td>
<td>Cumbersome</td>
<td>Easier</td>
</tr>
<tr>
<td>Limitations</td>
<td>Visibility and air contamination</td>
<td>Dose related toxicity and allergy</td>
</tr>
<tr>
<td>Delivery via small caliber catheters</td>
<td>Possible</td>
<td>Difficult</td>
</tr>
<tr>
<td>Radiation exposure</td>
<td>Increased if digital subtraction angiography used</td>
<td>Standard</td>
</tr>
<tr>
<td>Dose</td>
<td>Rate related toxicity</td>
<td>Volume related nephrotoxicity</td>
</tr>
<tr>
<td>Contraindications</td>
<td>Pulmonary-systemic communications; not for use in heart, brain or spinal vasculature</td>
<td>Allergy, nephrotoxicity</td>
</tr>
<tr>
<td>Hepatotoxicity</td>
<td>Rare</td>
<td>Rare</td>
</tr>
<tr>
<td>Quality of image</td>
<td>Good</td>
<td>Better</td>
</tr>
<tr>
<td>Procedure duration</td>
<td>Increased</td>
<td>Standard</td>
</tr>
</tbody>
</table>
as well as these are the presence of cardiac septal defects, pulmonary AV malformations, pulmonary hypertension and severe emphysema. In a series of 146 arteriovenous fistulography procedures, in 3 cases when manual injection of CO₂ into the brachial artery was performed, a reflux of the gas into the thoracic aorta occurred precipitating transient loss of consciousness[35].

Typically, CO₂ angiography does not cause any significant changes in the serum osmolality or blood gas values[40] unless excessive quantities of CO₂ are used or significant derangements of pulmonary function happen. Caution is required in cases where pulmonary functions are compromised such as in chronic obstructive pulmonary disease, as clearance of CO₂ may be decreased. Doses of CO₂ for diagnostic purposes are typically between 20-40 cc and it has no effect on ECG, and if possible capnography) need to be monitored.

Peristaltic and breathing movements sometime may decrease the image quality of mesenteric CO₂ angiography. This problem may be avoided by selective or superselective CO₂ injection into the mesenteric arterial branch, getting additional mask images or using intravenous glucagon to suppress the peristalsis. While using CO₂, sedation should be avoided or minimized as any of the side effects of CO₂ overdosing or air contamination may be missed in the presence of heavy sedation. During the procedure patients vital signs (pulsoximetry, blood pressure, heart rate, respiratory rate, ECG, and if possible capnography) need to be monitored closely. Any change in these parameters should raise the suspicion of CO₂ overdose or air contamination.

The utility of CO₂ as contrast agent for CT angiography for abdominal aorta and peripheral vessels is also currently being evaluated[41,42]. In an animal study the use of CO₂ micro-bubbles mixed in saline was compared with conventional CO₂ gas and ICA and demonstrated that vessels can be depicted using X-ray angiography and CO₂ micro-bubbles as enhancement[43]. CO₂ bubbles sometimes may provide better visualization then plain CO₂ gas with additional benefit of low dose requirement[44].

CONCLUSION
CO₂ is useful in cases where ICA cannot be used due to allergy or impaired kidney functions. CO₂ may be superior to ICA in certain procedures such as in TIPS.

REFERENCES
Ali F et al. Carbon dioxide as intravascular contrast agent


27 Wong AA, Charabel RA, Louie JD, Sze DY. Carbon dioxide contrast enhancement for C-arm CT utility for treatment planning during hepatic embolization procedures. *J Vasc Interv Radiol* 2013; 24: 975-980 [PMID: 23796085 DOI: 10.1016/j.jvir.2013.03.029]


P- Reviewer: Landesberg G, Maruyama H  S- Editor: Kong JX  L- Editor: A  E- Editor: Lu YJ