Differentiation of premacular hemorrhages with niveau formation

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Dear Editor,

We present three anatomical forms of premacular hemorrhage with niveau formation: hemorrhage in posterior precortical vitreous pocket (intrapocket hemorrhage), sub-internal limiting membrane (ILM) hemorrhage, and premacular retrocortical hemorrhage. Niveau formation is the formation of a horizonal boundary between fluid and blood cells when they accumulate in a closed space, because blood cells with higher specific gravity settle in a standing position. Differentiating between the three forms of hemorrhage is important for understanding the pathophysiology of premacular hemorrhage and selecting treatment.

Premacular hemorrhage may cause sudden loss of vision when the accumulation of blood is localized in the macular region. Various causes have been reported, such as retinal arterial macroaneurysm, Valsalva retinopathy, trauma, Terson syndrome, proliferative diabetic retinopathy, retinal vein occlusion, Eales’ disease, pregnancy and hematologic disorders[1-7]. Different treatment approaches have been used, including observation[1], neodymium-doped yttrium aluminum garnet (Nd:YAG) laser hyaloidotomy or membranotomy[2-5], and vitrectomy[6-7], but no clear guidelines are available. One reason is that the different anatomical forms of premacular hemorrhage are not differentiated before treatment; namely, hemorrhage in the posterior precortical vitreous pocket, hemorrhage behind the ILM, and hemorrhage between the ILM and the posterior vitreous cortex.

The vitreous of the human eye has a physiological liquefaction cavity before the macula. Worst[8] observed the vitreous by injecting Indian ink into an autopsy eye without retina, choroid or sclera. He reported for the first time that there was a space called the “bursa premacularis” in front of the macula. Kishi and Shimizu[9] stained the gel component of autopsy eyes with fluorescein and showed a space in front of the macula without posterior vitreous detachment (PVD), which was attached to the macula. They called this structure “posterior precortical vitreous pocket” (vitreous pocket) to differentiate it from bursa premacularis. From this anatomical relationship, it became possible to explain the pathogenic mechanism of vitreous macular diseases[10-11]. When triamcinolone was injected into the vitreous pocket during vitreous surgery, a disc-shaped glossy surface was observed, and the vitreous pocket was described as a liquefied lacuna surrounded by dense networks of fibrils[12]. Recently, a combination of ultra-widefield swept-source optical coherence tomography (OCT) and 3-dimensional structural imaging has been reported to be useful in visualizing the complex structure of the vitreous. These technologies are powerful tools that can be used to clarify the normal evolution of the vitreous, pathological changes of the vitreous, and implications of vitreous changes in various vitreoretinal diseases[13].

This study was conducted in accordance with the ethical principles of Declaration of Helsinki. Written informed consent for publication of patient information and images was given by the patients. The study was approved by the Ethics Committee of the Nihon University School of Medicine (No.161001).

CASE 1
A 50-year-old man diagnosed with Valsalva retinopathy had best corrected visual acuity (BCVA) of 20/400. Examination revealed a hemispherical hemorrhage 6 mm in diameter within the vitreous pocket centered over the macula. The pocket was surrounded by fibril networks, giving the surface of the intrapocket hemorrhage a shiny appearance. Bleeding in the pocket showed niveau formation. Since the vitreous pocket is almost a closed space, no spreading of bleeding to surrounding tissue was found, except for the hemorrhage in the connecting channel leading to Cloquet’s canal (Figure 1A). On OCT, no
retinal compression was seen as the hemorrhage was on the retina. Because of the thin anterior wall of the vitreous pocket (Figure 1B), the hemorrhage was denser in the inferior portion due to gravity. A space corresponding to the vitreous pocket posterior wall was visible. Four days later, the intrapocket hemorrhage was displaced inferiorly (Figure 1C) and the hemorrhage was absorbed spontaneously following formation of PVD (Figure 1D), and BCV A improved to 20/16.

CASE 2
A 47-year-old healthy female presented with decreased BCV A to 20/25 in her left eye associated with Valsalva retinopathy. Examination revealed a hemispherical hemorrhage beneath the ILM with a shiny surface showing niveau formation (Figure 2A). PVD was apparently absent on OCT. Since the ILM is not a closed structure, spreading of bleeding to surrounding tissue was observed (Figure 2B). On OCT, the sub-ILM hemorrhage was found to compress the retina, and the hemorrhage had uniform density throughout the whole depth because bleeding was beneath the ILM. The hemorrhage was absorbed after 3mo (Figure 2C), but BCV A remained unchanged at 20/25 due to cells depositing in the sub-ILM space of the fovea as shown on OCT (Figure 2D).

CASE 3
A 52-year-old man with premacular retrocortical hemorrhage associated with proliferative diabetic retinopathy presented with left BCV A of 20/400. Examination showed a hemispherical hemorrhage with niveau formation. Since the retrocortical space is not a closed structure, spreading of bleeding to surrounding tissue was observed (Figure 3A). Bleeding spread nasally beyond the optic disc, accompanied by vitreous hemorrhage. On OCT, the retina was not compressed by the premacular retrocortical hemorrhage (Figure 3B). Vitreous surgery and cataract surgery were performed, and retinal compression was seen as the hemorrhage was on the retina. Because of the thin anterior wall of the vitreous pocket (Figure 1B), the hemorrhage was denser in the inferior portion due to gravity. A space corresponding to the vitreous pocket posterior wall was visible. Four days later, the intrapocket hemorrhage was displaced inferiorly (Figure 1C) and the hemorrhage was absorbed spontaneously following formation of PVD (Figure 1D), and BCV A improved to 20/16.
The vitreous pocket in diameter and centered on the macula, coinciding with pocket. In intrapocket hemorrhage, bleeding occurs in the vitreous forms of premacular hemorrhage with niveau formation.

Table 1 summaries the differentiation between the three (Figure 3D).

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Intrapocket hemorrhage</th>
<th>Sub-ILM hemorrhage</th>
<th>Premacular retrocortical hemorrhage</th>
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</thead>
<tbody>
<tr>
<td>Site of hemorrhage</td>
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<td>Between ILM and posterior vitreous cortex</td>
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<tr>
<td>PVD Absent</td>
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<td></td>
<td></td>
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<tr>
<td>Fundus findings</td>
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<td>Hemispheric hemorrhage not centered on the macula</td>
<td>Irregular or hemispherical bleeding not centered on the macula</td>
</tr>
<tr>
<td>Size of vitreous pocket: about 6 mm in diameter</td>
<td>Various sizes: 1/4 to several disc diameters</td>
<td>Bleeding spreading to surrounding (+)</td>
<td>Various sizes: 1/4 to several disc diameters</td>
</tr>
<tr>
<td>Smooth surface (+)</td>
<td></td>
<td>Smooth surface (+)</td>
<td>Bleeding spreading to surrounding (+)</td>
</tr>
<tr>
<td>OCT findings</td>
<td>Denser lower portion by gravity due to the thin vitreous pocket anterior wall</td>
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<td>Denser at the site of PVD</td>
</tr>
<tr>
<td>Bleeding compresses retina (-)</td>
<td></td>
<td>Bleeding compresses the retina (+)</td>
<td>Bleeding compresses retina (-)</td>
</tr>
</tbody>
</table>

OCT: Optical coherence tomography; ILM: Internal limiting membrane; PVD: Posterior vitreous detachment.

Photocoagulation was also used (Figure 3C). After 5mo, BCVA recovered to 20/32, and OCT showed no macular edema (Figure 3D).

Table 1 summaries the differentiation between the three forms of premacular hemorrhage with niveau formation. In intrapocket hemorrhage, bleeding occurs in the vitreous pocket[9]; therefore, the hemorrhage is approximately 6 mm in diameter and centered on the macula, coinciding with the vitreous pocket[12]. Since bleeding occurs in the vitreous pocket, the retina is not compressed by the hemorrhage, and the space corresponding to vitreous pocket posterior wall can be seen. Intrapocket hemorrhage is recognized as a large hemispherical hemorrhage with a shiny surface accompanied by niveau formation. This form of hemorrhage occurs in association with conditions such as Valsalva retinopathy in the absence of PVD. Since the vitreous pocket is almost a closed space, no spreading of bleeding to surrounding tissue was found, except for the hemorrhage in the connecting channel leading to Cloquet’s canal. Because bleeding occurs below the thin anterior wall of the vitreous pocket, the hemorrhage is denser in the inferior region due to gravity.

The occurrence of sub-ILM hemorrhage is unrelated to the status of PVD. The size of hemorrhage ranges from 1/4 to several disc diameters. Large hemispherical hemorrhages show a shiny surface with niveau formation, but the location and size of the hemorrhage do not coincide with the vitreous pocket. As the ILM is not a closed structure, bleeding spreads slightly to surrounding tissue. Because bleeding occurs behind the rigid ILM, there is no density gradient in the hemorrhage due to gravity. Furthermore, since the ILM is a part of the retinal tissue, compression of the retina by the hemorrhage is clearly depicted in the peripheral region. Hemorrhage may be associated with subretinal or intraretinal hemorrhage.

Premacular retrocortical hemorrhage is caused by bleeding between the posterior vitreous cortex and ILM, associated with conditions such as proliferative diabetic retinopathy, in the presence of partial PVD. Although this hemorrhage may show niveau formation, bleeding spreads nasally beyond the optic disc, accompanied by vitreous hemorrhage. The hemorrhage has a flat irregular appearance and may spread irregularly, There is no compression of the retina by the hemorrhage.

Regarding treatment, although intrapocket hemorrhage may be absorbed spontaneously accompanying PVD formation, YAG laser treatment is selected when spontaneous absorption does not occur. An observation period of around one month would be appropriate. For the treatment of sub-ILM hemorrhage, observation, YAG laser and vitrectomy are used[1-7]. Waiting for spontaneous absorption is recommended for small hemorrhage. For large hemorrhages, many cases have been successfully treated by YAG laser, but multiple retinal holes after treatment have been reported[4]. Large hemorrhage may lead to proliferative vitreoretinopathy-like changes on the inner retinal surface of the ILM if surgery is delayed[10]. Many cases require treatment with anti-vascular endothelial growth factor therapy, vitrectomy, or photocoagulation for underlying diseases such as proliferative diabetic retinopathy.

Differentiation between intrapocket hemorrhage, sub-ILM hemorrhage, and premacular retrocortical hemorrhage by fundus photography and OCT is important for understanding the pathophysiology of premacular hemorrhages, which may help decide treatment approach. In this short report, we have clearly demonstrated the features of the three forms of macular hemorrhage in three cases. However, in clinical practice, differentiation of the premacular hemorrhages may not be straightforward due to morphological variations and differences in clinical background. More detailed clinical and imaging studies on a larger number of cases are required, and the effects on retinal tissues should be examined.

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REFERENCES