



# PREFERRED PRACTICE PATTERN® GUIDELINES

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## PREFERRED PRACTICE PATTERN®

- It is a committee developed by a panel of ophthalmologists with expertise in the guideline topic, a methodologist, and other experts.

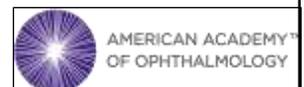


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American Society of Retina Specialists  
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 Central American Retina and Vitreous Society  
 European Society of Retina Specialists  
 The Macula Society  
 National Eye Institute  
 National Medical Association  
 Pan-American Retina and Vitreous Society  
 The Retina Society  
 Thai Retina Society

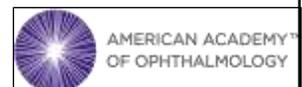


## PREFERRED PRACTICE PATTERN®

- For each PPP, a detailed literature search of PubMed and the Cochrane Library for articles in the English language is conducted
- The results are reviewed by an expert panel and used to prepare the recommendations, which are then given a rating that shows the strength of evidence when sufficient evidence exists.

## PREFERRED PRACTICE PATTERN®

- They provide guidance for practice not for the care of a particular individual.
- Preferred Practice Pattern® guidelines are not medical standards to be adhered to in all individual situations.
- They are reviewed annually and valid for 5 years.



## LEVELS OF EVIDENCE



## PREFERRED PRACTICE PATTERN® CLINICAL QUESTIONS

➤ PPP Clinical Questions answer the **PICTOS** questions:

- **P**atient.
- **I**ntervention.
- **C**omparison.
- **O**utcome.
- **T**ime frame.
- **S**tudy design or setting.

## SCOTTISH INTERCOLLEGIATE GUIDELINE NETWORK (SIGN)

|              |   |
|--------------|---|
| <b>I ++</b>  | <ul style="list-style-type: none"> <li>• High quality meta-analyses, systematic reviews of RCTs</li> <li>• RCTs with a very low risk of bias</li> </ul>   |
| <b>I +</b>   | <ul style="list-style-type: none"> <li>• Well-conducted meta-analyses, systematic reviews of RCTs</li> <li>• RCTs with a low risk of bias</li> </ul>  |
| <b>I -</b>   | <ul style="list-style-type: none"> <li>• Meta-analyses, systematic reviews of RCTs</li> <li>• RCTs with a high risk of bias</li> </ul>  |
| <b>II ++</b> | <ul style="list-style-type: none"> <li>• High-quality systematic reviews of case-control or cohort studies</li> <li>• High-quality case-control or cohort studies with a very low risk of confounding or bias and a high probability that the relationship is causal</li> </ul> |
| <b>II +</b>  | <ul style="list-style-type: none"> <li>• Well-conducted case-control or cohort studies with a low risk of confounding or bias and a moderate probability that the relationship is causal</li> </ul>   |
| <b>II -</b>  | <ul style="list-style-type: none"> <li>• Case-control or cohort studies with a high risk of confounding or bias and a significant risk that the relationship is not causal.</li> </ul>  |
| <b>III</b>   | <ul style="list-style-type: none"> <li>• Case reports and case series.</li> </ul>   |

## GRADING OF RECOMMENDATIONS ASSESSMENT, DEVELOPMENT AND EVALUATION (GRADE)

The body of evidence quality ratings are defined by GRADE as follows:

**Good quality (GQ)** • Further research is very unlikely to change our confidence.

**Moderate quality (MQ)** • Further research is likely to have an important impact on our confidence in the estimate of the effect.

**Insufficient quality (IQ)** • Further research is very likely to have an important impact on our confidence in the estimate of effect  
• Estimate of effect is very uncertain.

## GRADING OF THE IMPORTANCE IN THE CARE PROCESS

### Strong recommendation (SR)

- The desirable effects of an intervention **clearly** outweigh the undesirable effects or clearly do not

### Discretionary recommendation (DR)

- The trade-offs are less certain – either because of low-quality evidence or because evidence suggests that desirable and undesirable effects are closely balanced

## EXAMPLES

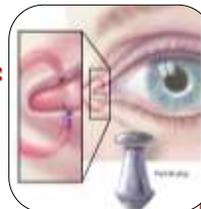
### Primary Angle Closure (Initial Evaluation and Therapy);

- Perform prophylactic iridotomy in fellow eye if chamber angle is anatomically narrow, as nearly half of fellow eyes can develop AACC within 5 years (II++, GQ, SR)



### Dry Eye Syndrome (Management Recommendations):

- For moderate dry eye, punctal plugs. (I++, GQ, SR)



### Age-Related Macular Degeneration (Initial and Follow-up Evaluation)

- Encourage patients who are currently smoking to quit. (I++, GQ, SR)





PREFERRED PRACTICE PATTERN®

**Diabetic Retinopathy**




**AMERICAN ACADEMY OF OPHTHALMOLOGY**  
*The Eye M.D. Association*

## HIGHLIGHTS AND RECOMMENDATIONS

Type 1 diabetes should have annual screenings for diabetic retinopathy beginning 5 years after the onset of their disease.

Type 2 diabetes should have a prompt examination at the time of diagnosis and at least yearly examinations thereafter. **II+; Good; Strong**

Patients should be informed of the importance of maintaining good **A1C** levels, serum lipids, and blood pressure to lower the risk of retinopathy developing and/or progressing. **II+; Good; Strong**

Patients with diabetes may use aspirin for other medical indications. **I+; Good; Discretionary**

Gestational diabetes do not require an eye examination. However, diabetic pregnant women should be examined early in the course of the pregnancy. **II+; Good; Strong**

At this time, laser photocoagulation remains the preferred treatment for non-center-involving diabetic macular edema. **I+; Good; Strong**

**TABLE 1 DIABETIC RETINOPATHY DISEASE SEVERITY SCALE AND INTERNATIONAL CLINICAL DIABETIC RETINOPATHY DISEASE SEVERITY SCALE**

| Disease Severity Level       | Findings Observable upon Dilated Ophthalmoscopy  |
|------------------------------|--|
| No apparent retinopathy      | No abnormalities   |
| Mild NPDR (see Glossary)     | Microaneurysms only  |
| Moderate NPDR (see Glossary) | More than just microaneurysms but less than severe NPDR  |
| Severe NPDR                  |  |
| U.S. Definition              | Any of the following (4-2-1 rule) and no signs of proliferative retinopathy: <ul style="list-style-type: none"> <li>• Severe intraretinal hemorrhages and microaneurysms in each of <b>four</b> quadrants</li> <li>• Definite venous beading in <b>two</b> or more quadrants</li> <li>• Moderate IRMA in <b>one</b> or more quadrants</li> </ul> |
| International Definition     | Any of the following and no signs of proliferative retinopathy: <ul style="list-style-type: none"> <li>• More than 20 intraretinal hemorrhages in each of <b>four</b> quadrants</li> <li>• Definite venous beading in <b>two</b> or more quadrants</li> <li>• Prominent IRMA in <b>one</b> or more quadrants</li> </ul>                          |
| PDR                          | One or both of the following: <ul style="list-style-type: none"> <li>• Neovascularization</li> <li>• Vitreous/previtreal hemorrhage</li> </ul>   |

## RECOMMENDATIONS

**TABLE 3 RECOMMENDED EYE EXAMINATIONS FOR PATIENTS WITH DIABETES MELLITUS AND NO DIABETIC RETINOPATHY**

| Diabetes Type                    | Recommended Initial Evaluation   | Recommended Follow-up*   |
|----------------------------------|--|--|
| Type 1                           | 5 years after diagnosis <sup>29</sup>                                    | Yearly <sup>29</sup>   |
| Type 2                           | At time of diagnosis <sup>31, 32</sup>                                   | Yearly <sup>31, 32</sup>   |
| Pregnancy†<br>(Type 1 or Type 2) | Soon after conception and early in the first trimester <sup>33, 34</sup> | <ul style="list-style-type: none"> <li>• No retinopathy to mild or moderate NPDR: every 3–12 months<sup>33, 35</sup></li> <li>• Severe NPDR or worse: every 1–3 months<sup>33, 36</sup></li> </ul> |

TABLE 6 MANAGEMENT RECOMMENDATIONS FOR PATIENTS WITH DIABETES

| Severity of Retinopathy | Presence of Macular Edema | Follow-up (Months) | Preretinal Photocoagulation (Scatter) Laser | Focal and/or Grid Laser* | Intravitreal Anti-VEGF Therapy |
|-------------------------|---------------------------|--------------------|---|--------------------------|--------------------------------|
| Normal or minimal NPDR  | No                        | 12                 | No  | No                       | No                             |
| Mild NPDR               | No                        | 12                 | No  | No                       | No                             |
|                         | ME<br>CSME†               | 4-6<br>1*          | No<br>No                                    | Sometimes                | Sometimes                      |
| Moderate NPDR           | No                        | 12†                | No  | No                       | No                             |
|                         | ME<br>CSME†               | 3-6<br>1*          | No<br>No                                    | Sometimes                | Sometimes                      |

Severe NPDR

**High-Risk PDR**

The presence of any three of the following four features characterizes DRS high-risk PDR:

- ◆ Neovascularization (at any location)
- ◆ Neovascularization at the optic disc
- ◆ Severe neovascularization:
  - New vessels within one disc diameter of the optic nerve head that are larger than one-quarter to one-third disc area in size
  - New vessels elsewhere that are at least one-half disc area in size
- ◆ Vitreous or preretinal hemorrhage

Non-high-risk PDR

High-risk PDR

The initial physical examination should include slit-lamp biomicroscopy: **III; Good; Strong**

The initial physical examination should include intraocular pressure: **III; Good; Strong**

The initial physical examination should include gonioscopy before dilation, when indicated: **III; Good; Strong**

## POSSIBLE APPLICATION IN EGYPT



### CARE PROCESS

It is recommended that an HbA1c of 7.0% or lower is the target for glycemic control in most patients while in selected patients, there may be some benefit to setting a target of 6.5%: **I++; Good; Strong**



## CARE PROCESS

Aspirin appears to be neither helpful nor harmful in the management of diabetic retinopathy: **I++; Good; Discretionary**



Focal, grid laser and anti-VEGF treatment sometimes recommended for patients with mild and moderate NPDR and CSME: **I++; Good; Strong.**



Anti-VEGF therapy is the treatment of choice for macular edema with or without focal laser treatment: **I++; Good; Strong.**

PRP treatment recommended for patients with DR starting from severe NPDR without DME and should not be delayed: **I++; Good; Strong.**



PROTOCOL 5

## CARE PROCESS

Many retina specialists prefer a less intense laser treatment, greater spacing, directly targeting microaneurysms, and avoiding foveal vasculature within at least 500  $\mu\text{m}$  of the center of the macula: **I++; Good; Discretionary**



PRP SHOULD NOT be recommended for eyes with mild or moderate NPDR, provided that follow-up [CAN] be maintained: **I++; Good; Strong**



The risk of severe visual loss among patients with high-risk PDR is reduced substantially by treatment using PRP as described in the DRS and ETDRS: **I++; Good; Strong**



Previously untreated PDR patients who have vitreous opacities and active neovascularization or fibrovascular proliferation should be considered to PPV: **I++; Good; Strong**



## SUMMARY

- **Preferred Practice Pattern**<sup>®</sup> is the highest level in the evidence pyramid.
- **Preferred Practice Pattern**<sup>®</sup> provide **guidance for practice** not for the care of a particular individual.
- **Preferred Practice Pattern**<sup>®</sup> guidelines are not medical standards to be adhered to in all individual situations.



**THANK  
YOU**