

High Sensitivity C-Reactive Protein: Emerging Biomarker for Primary Prevention of Cardiovascular Disease

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Increased CRP as measured using a high-sensitivity assay (hsCRP) is considered to be a very promising novel biochemical marker for the prediction of future coronary events. Several prospective studies have also demonstrated that hsCRP is a predictor of future cardiovascular morbidity and mortality among individuals with known cardiovascular disease.^{1,2} Recently completed study, clinical trial JUPITER³ has demonstrated that statins therapy to those with elevated hsCRP can significantly reduce future Cardiovascular disease (CVD). US National guidelines⁴ suggested that patients who are at intermediate risk (10%–20% 10-year predicted risk) for future cardiovascular events and who also have increased hsCRP (>3 mg/L) be considered for more aggressive vascular disease prevention strategies.

We had read with interest on recent reports of Ghosh et al "Prognostic value of baseline high-sensitivity C-reactive protein in patients undergoing replacement arthroplasty" "published in Journal of Nepal Medical Association 2009 vol 48 No 3 issue 174 page 144-8." It is very interesting to know elevated baseline hsCRP can predict the complication after post operative D14 of replacement arthroplasty. However, use of term high sensitivity on the article is not justifiable. High sensitivity refers detection of minimally elevated CRP and the technique usually have detection limit less than 0.03 mg/dL (0.3 mg/L). CRP greater than 1.0 mg/dL can be routine estimated in laboratory and doesn't refer high sensitivity. Recent laboratory medicine guideline by National academy of clinical biochemistry,⁵ recommended that hsCRP results, regardless of the method used, should be expressed as mg/L. Further, hsCRP assays categorizes patients as Low risk <1.0

mg/L, Average risk 1.0–3.0 mg/L, High risk <3.0 mg/L, Very high risk ≥ 10.0 mg/L. US National guidelines (2003) also suggest similar categories but it had recommended hsCRP ≥ 10.0 mg/L as indicator of other inflammatory diseases. Ghosh et al had used unit of mg/dL and categorized patients on the basis of hsCRP as <3.0 mg/dL and >3.0 mg/dL that is (<30.0 mg/L and > 30.0mg/L). Such highly elevated condition can be seen only in the state of inflammation and are detected by routine CRP not by so called hsCRP.

REFERENCES

1. Rifai N, Ridker PM. High-sensitivity C-reactive protein: a novel and promising marker of coronary heart disease. Clin Chem. 2001;47:403-11.
2. Ridker PM. C-Reactive Protein: Eighty Years from Discovery to Emergence as a Major Risk Marker for Cardiovascular Disease. Clin Chem. 2009;55(2):209-15.
3. Ridker PM, Danielson E, Fonseca FAH, Genest J, Gotto AM, Kastelein JP et al. On behalf of the JUPITER Trial Study. Rosuvastatin for vascular prevention among men and women with elevated C-reactive protein. N Engl J Med. 2008;359:2195-207.
4. Pearson TA, Mensah GA, Alexander RW, Anderson JL, Cannon RO, Criqui M, et al. Markers of inflammation and cardiovascular disease: application to clinical and public health practice: a statement for healthcare professionals from the Centers for Disease Control and Prevention and the American Heart Association. Circulation. 2003;107:499-511.
5. Myers GL, Christenson RH, Cushman M, Ballantyne CM, Cooper GR, Pfeiffer CM, et al. National Academy of Clinical Biochemistry Laboratory Medicine Practice Guidelines: Emerging Biomarkers for Primary Prevention of Cardiovascular Disease. Clin Chem. 2009;55:378-84.

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