C-Reactive Protein in Guillainbarré Syndrome: Recent Advances and Future Prospective

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Abstract

Introduction: C-reactive protein (CRP) is an acute phase reactant (APR), which is widely used as a potential biomarker in various infectious and inflammatory conditions (IICs). C-reactive protein (CRP) is also responsible for the Guillain-Barré syndrome (GBS).

Aim and Objectives: C-reactive protein (CRP) levels were estimated in sera of patients with GBS and compared with adequate healthy controls from 2014 to 2018 from our locality.

Settings and Design: This is a prospective-control study from (2014-2018) from our locality.

Study Population: This study divided into 4 groups: 1. GBS group included 50 newly diagnosed patients with GBS; 2. Neurological control (NC) group comprised of 50 patients with non-paralytic neurological symptoms/disorders (NNPS); 3. Non-neurological controls (NNC) comprised of 50 patients having no neurological symptoms and 4. Healthy controls group (HCG) comprised of 150 healthy subjects.

Materials and Methods: C-reactive protein (CRP) was evaluated by slide latex agglutination test (SLAT) and enzyme linked immunosorbent assay (ELISA) through available kits in the market or available from the noted company.

Statistical Analysis: We used the very simple and descriptive statistics through available software in SPSS Statistics versions 16.0.

Results: CRP by LAT was positive in 25.8% GBS group, 35.5% NC group and 45.6% NNC group. The range of titer in CRP positive samples in the three patient groups (GBS, NC, NNC) was at concentration of 0.5 mg/dl to 19.5 mg/dl. The similar results were also obtained by ELISA in the patient groups (PGs). None of the HC subjects was positive for detectable levels of CRP were observed. Additionally, the Higher basal level (HBL) of CRP was detected in patients with GBS successfully.

Conclusion: Autoimmune conditions like Guillain-Barré syndrome (GBS) can stimulate the production of a higher level of inflammation rate (HILR) resulted through C-reactive protein (CRP) is a unique biomarker for the Bilateral facial nerve palsy (BFNP) or facial diplegia (FD) is a rare condition that occurs mainly in the context of Guillain-Barré syndrome (GBS).

Keywords: C-reactive protein; Guillain-Barré syndrome; Slide latex agglutination test; Enzyme linked immunosorbent assay; Violent systemic autoimmune reaction

Abbreviations: CRP: C-Reactive Protein; IICs: Infectious and Inflammatory Conditions; APR: Acute Phase Reactant; GBS: Guillain-Barré Syndrome; HIPM: Humoral Immuno Pathogenic Mechanism; AAIM: Acute Autoimmune Disorder; HBL: Higher Basal Level; ALTs: Autoimmunological Titters; PLR: Prolonged Leukemoid Reaction; RF: Rheumatoid Factor; ANCA: Antineutrophil Cytoplasmic Autoantibody; ARDS: Acute Respiratory Distress Syndrome; MS: Multiple Sclerosis; MI: Myocardial Ischemia; COPD: Chronic Obstructive Pulmonary Disease; TD: Tissue detection; TAMA: Therapeutic Anti-Inflammatory Agent; DOMs: Dissemination of Microorganisms; FI: Focal Infection; EC: Eudro-constitutionality; ALS: Amyotrophic Lateral Sclerosis

Introduction

C-reactive protein (CRP) is an acute phase reactant (APR), used as a biomarker for various infectious and inflammatory conditions (IICs) [1]. Guillain-Barré syndrome (GBS) is an acute autoimmune, polyradiculoneuropathy (PRNP) triggered by infectious agents such as Campylobacter jejuni [2]. GBS is generally precipitated with 1-3 weeks following C. jejuni infection and suggests a humoral immunopathogenic mechanism (HIPM) [3]. The incidence rates of pyrexia, headache, nausea, gastro interological (GI) problems with GBS is related to acute autoimmune disorder (AAID) still unknown [4]. CRP levels were estimated within these of patients with GBS and compared with adequate controls and range of titer in CRP positive samples [5]. The three major patient groups (GBS, NC, NNC) were at
Concentration of 0.6 mg/dl to 19.2 mg/dl. The higher basal level (HBL) of CRP was also detected in patients with GBS [6]. In blood plasma exchange had been given as severe immunological reaction (VIR) was unable to control the acute motor axonal neuropathy (AMAN) and GBS [5]. Prolonged leukemoid reaction (PLR) and higher levels of autoimmunological titers (AIT), including C-reactive protein (CRP), rheumatoid factor (RF), and antimyeloperoxidase autoantibody (ANCA) also persisted with GBS [7]. Patient developed with acute respiratory distress syndrome (ARDS) and type 3 of rapidly progressive glomerulonephritis (RPGN) rapidly declined of renal function (RF) and immune suppressants were still not documented [8]. AMAN with the violent systemic autoimmune reaction (VSAIR) is strongly associated with infectious disease GBS [9]. GBS patients were implied with standard plasmapheresis (SPF) is insufficient and corticosteroids with immunosuppressant added in early stage of GBS is still a research question with (AMAN) [10].

Materials and Methods

Aim and objectives

C-reactive protein (CRP) levels were estimated in sera of patients with GBS and compared with adequate healthy controls from 2014 to 2018 from our locality/local area and informed.

Settings and design

This is a prospective-control study from 2014-2018 from our locality in prospective way.

Study population

This study we divided into 4 groups: 1. GBS group included 50 newly diagnosed patients with GBS; 2. Neurological control (NC) group comprised of 50 patients with non-paralytic neurological symptoms/disorders (NNPS) 3. Non-neurological controls (NNC) comprised of 50 patients having no neurological symptoms and 4. Healthy controls group (HCP) comprised of 150 healthy subjects on prior order.

Inclusion criteria

Who is willing to the study subject were included in this study.

Exclusion criteria

Those unwilling to this study subject were strictly excluded from our study.

Materials and methods

C-reactive protein (CRP) was evaluated by slide latex agglutination test (SLAT) and enzyme linked immunosorbent assay (ELISA) through available kits in the market or available from the noted company according to International standard protocol.

Statistical analysis

We used the very simple and descriptive statistics through available software in SPSS Statistics versions 16.0. Available on online and down loaded freely available.

Results

According to our findings of CRP by LAT was positive in 25.8% GBS group, 35.5% NC group and 45.6% NNC group. The range of litter in CRP positive samples in the three patient groups (GBS, NC, NNC) was at concentration of 0.5 mg/dl to 19.5 mg/dl. The similar results were also obtained by ELISA in the patient groups (PGs). None of the healthy control (HC) subjects was positive for detectable levels of CRP were observed. Additionally, the Higher basal level (HBL) of CRP was detected in patients with GBS successfully is our one of the great achievements in this study groups. LAT was positive in 25.8% and its nearly related patients (PCs) was unable for detection of CRP. Also, the comparison of CRP was not documented.

Discussion

αB-crystallin (HspB5) is a chaperone and a potential biomarker of innate immunity activation (IIA) for GBS [11]. Therapeutic potential has been investigated in several inflammatory diseases (multiple sclerosis (MS), myocardial ischemia (MI), and Guillain-Barré syndrome (GBS) and obstructive pulmonary disease (COPD) pathogenesis and their inflammation [12]). Tissue detection (TD) of the protein showed significantly overexpressed in COPD smokers in comparison to COPD nonsmokers expressed in patients with age-related emphysema [13]. αB-crystallin is a regulator of innate immunity (II) and a therapeutic anti-inflammatory agent (TIA) [14]. Focal infection (FI), has been localized or generalized infection caused by the dissemination of microorganisms (DOMs) or toxic products from a focus of infection in organic districts [15]. An "immuno-allergic theory" with formation of auto-antibodies in human body, genesis of autoimmune illnesses (AII) sustained by individual reactivity (IR) linked to eredo-constitutionality (EC) [16]. The local origin pathology (FOP), are cancer, sarcoidosis, multiple sclerosis (MS), amyotrophic lateral sclerosis (ALS), autism and Guillain-Barré syndrome (GBS) [17]. An Autoimmune Neuropsychiatric Disorders (AINPDs) were associated with Streptococcal infections (PANDAS), Tourette's syndrome (TS), myasthenia gravis (MG), polycystic kidney disease (PKD), obesity, Alzheimer's disease (AD) and diabetes mellitus (DM) among pediatric test had performed through by the Laboratory test analyses (LTA) is one of the important and strictly priority basis test (PBT) [18].

Lescocytic fluid (LF), protein electrophoresis (PF), Gel analysis, C-reactive protein, REUMA test VES, TES, were the test of an inflammatory process (IP). The DNA-Polymerase Chain Reaction method (PCR) and Real time Polymerase Chain Reaction method (RT-PCR) is the fundamental test for the diagnosis of bacterial and viral infections (BVI) [19]. An intensive periodontal therapy (IPT) also showed a significant reduction of lymphocyte formula at CRP levels of interleukin-6 (IL-6), (IL-8), Polymorphisms of IL-17 and ICAM-1 and their expression and the LDL cholesterol after two-three months of GBS [20]. Fever and acute respiratory failure (FARF), progressive tetra paresis,
delirium (PTD), behavioral abnormalities (BA), and diplopia (21). At leukocytosis had increased the C-reactive protein (CRP), present in appropriate anti-diuretic hormone secretion (SIADH) (22). The Lumbar puncture (LP), yielded colorless CSF with mononuclear pleocytosis (MNP) and Regulatory T cell frequency, but not plasma IL-33 levels the protein rises in axonal degeneration (AD) (23). An Electrodiagnosis test revealed that demyelinating neuropathy (DMNP) and axonal degeneration (AD) (24). The serum IgG and IgM for mycoplasma pneumoniae (MP) was consistent with acute infection (AI), erythrocytosis has been started with rapid resolution (RR) of the symptoms (25). Respiratory disease (RD) were not present and SIADH patient without the direct central nervous system (DCNS) involvement, differential diagnosis (DD) and possible pathogenic mechanisms (PPM) involved in Guillain-Barré syndrome (GBS) (26).

Recent advances and future prospective
ST-segment elevation (STE) noted in association with reversible left ventricular dysfunction (LVD) due to the synergistic toxic effect (STE) (27). An increased catecholamine and transiently damaged sympathetic nerve (DSN) endings at the myocardium in Guillain-Barré syndrome (GBS) (28). Importantly, causative Hantavirus serotype (HVT), the degree of "systemic inflammatory response syndrome" (SIRS) responsible to determine the clinical picture in Hantavirus infections and associated diseases of GBS (29). GBS can stimulate the production of a higher level of inflammation (HILO), increase in the CRP production on Bilateral facial nerve palsy (BLFP) or facial diplegia (FD) distinctively. We report patients with isolated idiopathic bilateral facial nerve palsy (IIBFP), with meninges having to evidence of Guillain-Barré syndrome (GBS) and Comparative Value of Erythrocyte Sedimentation Rate (ESR) and C-Reactive Protein (CRP) rapid and complete recovery till date (30).

Conclusion
We concluded that an unusual variant of bilateral facial palsy (UVBFPS), Acute motor axonal neuropathy (AMAN), Guillain Barre syndrome (GBS), is frequently induced by the antecedent infection via atypical pathogen, such as Campylobacter jejuni, Mycoplasma pneumonia and some viruses generally accepted that corticosteroids and immunosuppressants are not recommended in patients with (GBS including AMAN). Autonomic conditions like Guillain-Barré syndrome (GBS) can stimulate the production of a higher level of inflammation rate (HILO) resulted through C-reactive protein (CRP). Bilateral facial nerve palsy (BFNP) or facial diplegia (FD) is a rare condition that occurs mainly in the context of Guillain-Barré syndrome (GBS) C-reactive protein count as a potential and capable biomarker for such kind of activities.

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References


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