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Vaccinations in autoimmune myasthenia gravis

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Chapter 5

Influenza vaccination in patients with myasthenia gravis

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1. INTRODUCTION

Myasthenia gravis (MG) is an acquired autoimmune disease of the neuromuscular junction and is characterized by fluctuating weakness and fatigability of skeletal muscles [1]. In the majority of MG patients acetylcholine receptor (AChR) antibodies are found [1]. Symptomatic treatment is often insufficient, and a considerable proportion of patients need long-term immunosuppressive medication (IM). Patients with an autoimmune disorder are generally believed to be at an increased risk of infection, either due to their immunosuppressive therapy or due to immune abnormalities associated with their disease [2, 3]. Conversely, an infection can cause exacerbation of symptoms, potentially resulting in a myasthenic crisis. Specific data on infection rates in myasthenic patients are not available [4].

Little is known about the efficacy and safety of vaccination in patients with autoimmune diseases. No specific guidelines regarding vaccinations in patients with MG exist, but a small number of observational studies suggest that influenza vaccination is safe [5-7] and recently a randomized controlled trial showed that influenza vaccination has no influence on AChR antibody titres [8]. In a recent study we found a small, temporary, but significant increase in Quantitative Myasthenia Gravis scores (QMG) after tetanus vaccination. However, this was far less than what is generally considered clinically relevant [9]. In the Netherlands, annual vaccination against influenza is recommended for all patients with an autoimmune disease [10]. However, in our personal experience and as described earlier [5], many patients express concern that vaccination may lead to an exacerbation and a substantial number decline vaccination each year based on these concerns. This is unfortunate, as seasonal vaccination against influenza is highly effective in reducing laboratory-confirmed influenza illness, hospital admissions and risk of death, especially in elderly and frail patients [11]. This is relevant, as this age group has the highest incidence of autoimmune MG [12]. Another concern is that IM may hamper the development of protective antibody levels. Therefore, we performed a double-blind placebo-controlled trial to investigate the efficacy and safety of the seasonal (2016/2017) influenza vaccine in patients with AChR MG with and without IM.

2. MATERIALS AND METHODS

2.1 Standard protocol approvals, registrations, and patient consents

This study was approved by the Local Committee on Medical Ethics of the LUMC. Subjects provided written informed consent and received reimbursement of travel costs. The trial is listed on clinicaltrialsregister.eu under 2016-003138-26.

2.2 Patients

We included 47 patients with AChR MG and 47 healthy controls at the start of the flu season (October 2016). AChR MG patients were recruited from the neurology outpatient clinic of Leiden University Medical Center (LUMC) and through the

national patient organization. Seasonal influenza vaccination was offered at the start of the flu season to all LUMC employees; healthy controls were recruited from this population.

Inclusion criteria for the patient group were a diagnosis of AChR MG, age ≥ 18 years and stable disease in the past 3 months. Diagnosis of AChR MG was based on clinical signs or symptoms consistent with MG and a positive serological test for AChR antibodies. A maximum daily dose of 30mg of prednisolone, with a variation of ± 5 mg during 3 months before participation was allowed as well as use of other immunosuppressive medication.

During the study, patients were on a stable dose of their medication (see Table 1). Time from last pyridostigmine dose to clinical testing was kept constant for each patient on test days, but was allowed to vary between patients. Inclusion criteria for healthy controls were an age ≥ 18 years and no autoimmune disease or immunosuppressive medication.

Exclusion criteria for the AChR MG group were: instable or severe disease as evidenced by recent changes in medication or an MGFA classification of 4 or 5, presence of a thymoma, use of vitamin K antagonist or new oral anti-coagulants (NOACs), pregnancy and other diseases of the immune system that may affect the efficacy of vaccination.

2.3 Study protocol

This single-center, prospective, double-blind, randomized, placebo-controlled study was performed at the LUMC. Randomization was performed by a randomization list created by the hospital pharmacy. Patients and physicians performing clinical tests were blinded for treatment allocation until the end of T1. Research nurses, who administered the vaccination, were not blinded, because the placebo was provided in a different syringe than the commercial influenza vaccine. Patients were randomized to receive either an intramuscular injection with the influenza vaccine or a placebo (0.5 mL 0.9% NaCl) (T0). At T0 age, sex, disease duration, use of medication, MGFA classification, thymectomy and seasonal influenza vaccinations in the previous 3 years were recorded. Prior to injection (T0) and four weeks later (T1), serum and several clinical outcome measures were obtained. Four weeks (T1) after this first vaccination, patients were unblinded and patients in the placebo group were vaccinated with the influenza vaccine (Figure 1). At T2, 4 weeks after the flu vaccination, a third blood sample and MG specific activities of daily living (MG-ADL) score were obtained from the (initial) placebo group. In all patients, an MG-ADL was obtained by phone by a research nurse, twelve weeks after influenza vaccination (T3). At T1, T2 and T3 AChR MG patients were asked for side effects and exacerbation of their MG symptoms. Healthy controls were asked for side effects at T1. Figure 1 shows an overview of the study design.

	AChR MG Vaccination	AChR MG Placebo	HC	Total
Number of patients - n	24	23	47	94
Gender, female (%)	11 (45.8)	14 (60.9)	36 (76.6)*	61 (64.9)
Age, median years (range)	61.5 (32-72)	63 (22-74)	54 (24-65)*	
Duration of disease, mean years (SD)	14.3 (13.9)	10.7 (9.9)	-	
MGFA classification**			-	
0 - n (%)	10 (41.7)	8 (34.8)	-	
1 - n (%)	0	5 (21.7)	-	
2 - n (%)	13 (54.2)	9 (39.1)	-	
3 - n (%)	1 (4.2)	1 (4.3)	-	
Use of immunosuppressive medication, n (%)	15 (62.5)	14 (60.9)	-	
Prednisolone, n (%)	9 (37.5)	11 (47.8)	-	
Mean daily dose, mg (range)	9.2 (5-20)	6.8 (1-10)	-	
Azathioprine, n (%)	13 (54.2)	10 (43.5)	-	
Mean daily dose, mg (range)	131.2 (50-200)	116.7 (50-200)	-	
Mycophenolic acid, n(%)	0	2 (8.7)	-	
Mean daily dose, mg (range)	-	2000 (2000)	-	
Cyclosporine, n (%)	3 (12.5)	0	-	
Mean daily dose, mg(range)	166.7 (150-200)	-	-	
Combination of immunosuppressive medication, n (%)	8 (33.3)	8 (34.8)	-	
Thymectomy in the past (>1 year ago) - n (%)	15 (62.5)	14 (60.9)	-	
Past seasonal trivalent inactivated influenza vaccination - n (%)			39 (83)	78 (83)
2015-2016	15 (62.5)	16 (69.6)	28 (59.6)	59 (63)
2014-2015	16 (66.7)	17 (73.9)	33 (70.2)	66 (70)
2013-2014	16 (66.7)	15 (65.2)	31 (66)	62 (66)

Table 1. Baseline characteristics. The AChR MG group is divided in the in vaccination and placebo group. *Healthy controls are significantly younger ($p=0.001$) than the AChR MG group en consist out of significantly more females ($p=0.02$). ** MGFA classification: Myasthenia gravis foundation America classification.

2.4 Influenza vaccine

We used the commercially available influenza vaccine manufactured by Sanofi Pasteur (Vaxigrip, RVG 22306) for the season 2016/2017. One dose of 0.5 mL contains 15 µg haemagglutinin of each of the influenza virus strains in the split inactivated influenza vaccine: A/California/7/2009 (H1N1)pdm09, A/Hong

Kong/4801/14 (H3N2) and B/Brisbane/060/08 (B/Victoria/2/87- line). The vaccine was administered intramuscularly, as a bolus, in the non-dominant upper arm.

2.5 Influenza antibody response

The primary endpoint of this study was change in titre of antibodies to the flu vaccine strains. A secondary endpoint was the effect of IM on the humoral response. Antibodies to the vaccine strains A/California/7/2009 (H1N1)pdm09, A/Hong Kong/4801/14 (H3N2) and B/Brisbane/060/08 were measured using the hemagglutination-inhibition (HI) assay, according to standard methods at the national influenza center at the Erasmus Medical Center[13]. Titres below the detection limit (i.e. $\leq 1:10$) were assigned a value of 1:5. Geometric mean titres (GMTs) and seroprotection rates (defined as HI titres $\geq 1:40$) were chosen as the main outcome measures. Seroconversion was defined as a post-vaccination HI titre of at least 1:40 combined with at least a four-fold increase in titre. A non-responder was defined as a post vaccination HI-titre of $< 1:40$.

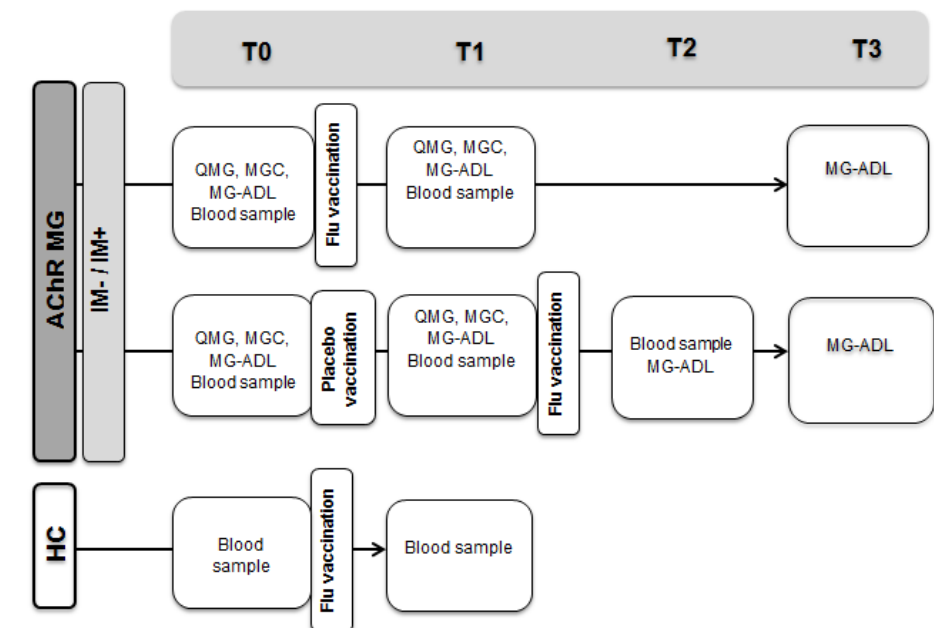


Figure 1. Study flowchart. T0: baseline, T1: 4 weeks after influenza (flu) or placebo vaccination, T2: 4 weeks after vaccination in placebo group, T3: 12 weeks after vaccination with influenza. In the AChR MG vaccination group a blood sample was taken at T0 and T1 and in the AChR MG placebo group at T0, T1 and T2 in the placebo group. HC: Healthy controls. QMG: Quantitative Myasthenia Gravis score, MGC: Myasthenia Gravis Composite score, MG-ADL: myasthenia gravis activities of daily living score.

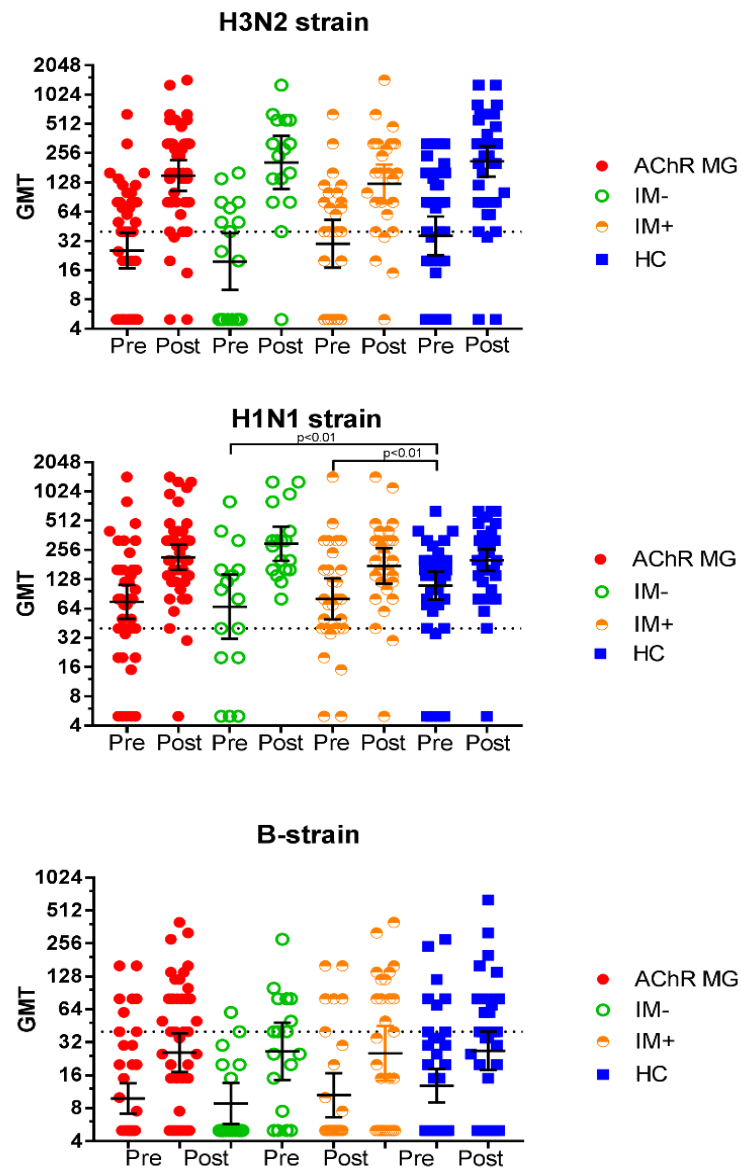


Figure 2. Response to influenza vaccination. Geomean titres (GMT) of H3N2, H1N1 and B-strain, pre and 4 weeks post-vaccination with a 95%CI. Groups consist of: 47 AChR MG patients, 18 AChR MG patients without immunosuppressive medication (IM-), 29 AChR MG patients with IM+ and 47 healthy controls. The dotted line is the minimal GMT that is considered as protective (HI-titre 1:40).

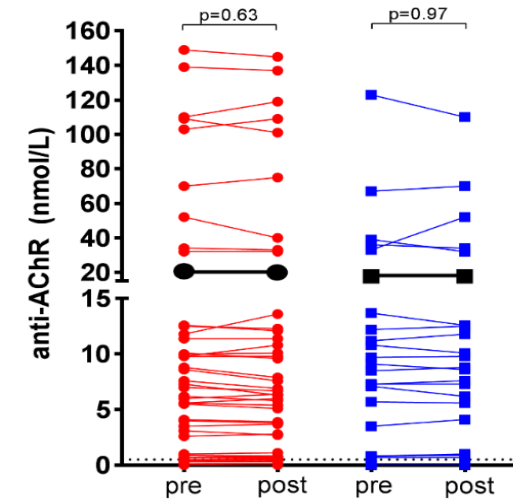


Figure 3. Anti-AChR antibody concentrations before and 4 weeks after vaccination with influenza in all MG patients and before and 4 weeks after placebo administration. The dotted line indicate the minimal titre that is considered as positive (0.5 nmol/L). Black: mean titre of the group, individual titres are depicted in colour (● Vaccination group (n=47); ■ Placebo group (n=23)).

2.6 Sampling protocol and clinical scoring

Another secondary endpoint was a clinically relevant change in clinical scores. We used the QMG, MG Composite (MGC) and the MG-ADL scores as clinical outcome measures. The QMG is a 13-item scale that measures muscle strength and endurance, ranging from 0 to 39. The MGC is a composite scale selected from existing MG-specific scales (MG-ADL, QMG and Manual Muscle Test (MMT)), ranging from 0 to 50. The MG-ADL is a scale to assess MG symptoms that patients experience in their daily activities, ranging from 0 to 24. A change of 2.3 points for the QMG, 3 for the MGC and 2 points for the MG-ADL was considered clinically relevant [14-16]. For all three outcome measures, higher scores indicate a higher clinical severity of MG [14-18].

2.7 Antibodies against AChR

The last secondary endpoint was a change in antibodies against AChR. AChR antibody titres were measured with a commercially available radio immunoprecipitation assay (RIA)(RSR Ltd., Cardiff, UK)[19]. Absolute titres were measured using multiple dilutions of each serum sample.

2.9 Statistical analysis and power

The study was powered for an expected response rate (i.e. seroprotection rates) of 75% with a 95%-confidence interval of 63-87% in MG patients. Therefore, 50 patients with MG were needed. Statistical analysis was performed with Graph-

Pad Prism software version 7 and SPSS version 23. In all tests $p < 0.05$ was considered statistically significant. Influenza titres were log transformed in order to normalize the data. Comparison for normally distributed numerical variables was done with paired or unpaired T-tests or a one-way analysis of variance (ANOVA). Influenza virus specific antibody responses were compared between AChR MG patients (with and without immunosuppression) and healthy controls. Within the AChR MG group, patients with and those without thymectomy (Tx) were compared. The AChR antibody titres before vaccination of all AChR MG patients were compared to titres 4 weeks after influenza vaccination. The clinical outcome measures were compared between the AChR MG vaccination and placebo group.

3. RESULTS

3.1 Patient characteristics

Forty-seven patients (53.2% female, median age 62 years, range 22-74 years) and 47 healthy controls (76.6% female, median age 54 years, range 24-66 years) were vaccinated with the seasonal influenza vaccine from October to December 2016. Healthy controls were significantly younger ($p = 0.001$) and were more frequently female ($p = 0.02$) than the MG group. In the MG group, 23 patients randomly received a placebo injection followed by flu vaccination 4 weeks later. Baseline characteristics did not differ between the two MG patients groups that either received first the flu vaccination or the placebo vaccination. The MG group consisted of 29 patients with (IM+) and 18 without (IM-) immunosuppressive medication. The IM+ group was significantly older ($p < 0.01$) than the IM- group and contained more female patients ($p = 0.04$). Disease duration and whether a patient underwent a thymectomy in the past was not significantly different between IM- and IM+ groups ($p = 0.4$ and $p = 0.16$, respectively). Baseline characteristics are given in Table 1.

3.2.1 Serological response to Influenza vaccination

Upon vaccination the MG group ($n = 47$) developed a geomean titre (GMT) for all three vaccine strains that was similar to the HC group (H3N2, $p = 0.2$; H1N1, $p = 0.7$; and B-strain, $p = 0.9$) (Figure 2). The post-vaccination seroprotection and seroconversion rates were comparable between the MG group and HC group for all strains. In the MG group, 40.4% of all patients (19/47) reached a seroprotective titre for all three strains. In the HC group this was 51% (24/47) (Table 2).

3.2.2 Influence of use of immunosuppressive medication and thymectomy

No significant effect on the serological response to influenza vaccination was observed between the IM- ($n = 18$) and IM+ group ($n = 29$) (H3N2, $p = 0.2$; H1N1, $p = 0.1$; and B-strain, $p = 0.9$). The pre-vaccination H1N1 GMT was significantly lower in both the IM- and IM+ groups ($p < 0.01$ for both), but there was no significant difference in post-vaccination GMT compared to the HC group. Seroconversion and post-vaccination seroprotection rates were also similar between HC and the IM- and IM+ groups (Table 2).

	AChR MG (n=47)	IM- (n=18)	IM+ (n=29)	HC (n=47)
H3N2 strain				
Pre HI titre $\geq 1:40$ - n (%)	25 (53.2)	8 (44.4)	17 (58.6)	26 (55.3)
Post HI titre $\geq 1:40$ - n (%)	42 (89.4)	17 (94.4)	25 (86.2)	44 (93.6)
Pre GMT - value (95% CI)	26 (17-39)	20 (10-39)	30 (17-53)	36 (23-57)
Post GMT - value (95% CI)	150 (104-216)	205 (109-384)	124 (78-196)	210 (147-301)
Seroconversion - n (%)	22 (46.8)	11 (61.1)	11 (37.9)	26 (55.3)
H1N1 strain				
Pre HI titre $\geq 1:40$ - n (%)	37 (78.7)	13 (72.2)	24 (82.7)	42 (89.4)
Post HI titre $\geq 1:40$ - n (%)	45 (95.7)	18 (100)	27 (93.1)	46 (97.9)
Pre GMT - value (95% CI)	75 (50-112)	67 (31-143)	80 (49-131)	110 (79-153)
Post GMT - value (95% CI)	215 (159-291)	297 (198-446)	176 (115-268)	201 (156-259)
Seroconversion - n (%)	15 (31.9)	6 (33.3)	9 (31)	9 (19.1)
B- strain				
Pre HI titre $\geq 1:40$ - n (%)	8 (17)	2 (11.1)	7 (24.1)	10 (21.3)
Post HI titre $\geq 1:40$ - n (%)	22 (46.8)	9 (50)	13 (44.8)	24 (51)
Pre GMT - value (95% CI)	10 (7-14)	9 (6-14)	11 (7-17)	13 (9-18)
Post GMT - value (95% CI)	26 (17-39)	26 (15-48)	25 (14-45)	27 (18-40)
Seroconversion - n (%)	12 (25.5)	7 (38.9)	5 (17.2)	13 (27.7)

Table 2. Humoral response to seasonal influenza vaccine 2016-2017. Chi-square tests showed no significant difference in pre and post HI titres between HC and AChR MG groups and between HC and IM-/IM+ groups.

Since the antibody response to influenza is T-cell dependent and a large portion of our patients (42.6%) underwent a thymectomy in the past (Table 1), we tested whether a thymectomy impacted the antibody response. We found no significant difference in pre- (H3N2, $p = 0.7$; H1N1, $p = 0.6$; B-strain, $p = 0.5$) and post-vaccination GMT (H3N2, $p = 0.2$; H1N1, $p = 0.4$; B-strain, $p = 0.5$), neither between patients with and without thymectomy, nor between patients and healthy controls (data not shown).

Both IM use and thymectomy can influence the absolute cell counts of T- and B-cells, therefore, we performed an immunophenotyping in all patients pre- and post-vaccination. Patients of the IM+ group had significantly lower absolute cell counts of CD19+ B-lymphocytes (mean $73 \times 10^6/L$, $p < 0.001$), CD4+ T-lymphocytes (mean $621 \times 10^6/L$, $p = 0.02$), CD8+ T-lymphocytes (mean $245 \times 10^6/L$, $p = 0.04$) and NK-cells (mean $97 \times 10^6/L$, $p < 0.001$) than patients of the IM- group. However, these values are in the range of healthy controls, except for the CD8+ T-lymphocytes

(normal values $260-990 \times 10^6/L$). There was no difference in absolute cell counts between the groups with and without thymectomy.

3.3 Non-responders

There were 5 non-responders in the MG group to H3N2 vaccination vs. 3 in the HC group, 2 to H1N1 vs. 1 in the HC group, 25 to the B-strain vs. 23 in the HC group. In the IM- group and IM+ group there were 1 and 4 non-responders respectively to H3N2, 0 and 2 respectively to H1N1, 9 and 16 respectively to the B- strain. The largest difference in response between IM- and IM+ groups was found for the B-strain: 9 non-responders in the IM- group and 16 in the IM+ group, although this apparent difference did not reach statistical significance: $p=0.73$. Of the 16 non-responders to the B-strain in the IM+ group, 12 used prednisone, 14 used azathioprine and 2 used three types of immunosuppressive medication (prednisone, azathioprine and cyclosporine). Only 1 MG patient and 1 HC were non-responders for all three strains.

3.4 Clinical scores

Figure 4 shows individual clinical scores and changes of the MG vaccination group ($n=24$) and MG placebo group ($n=23$) from T0 to T1. Use of IM was comparable (Table 1). Total scores of the three outcome measures were the same before and after vaccination between both groups. In addition, there was no significant change in the mean score or delta of all three outcome measures between T0 and T1. The MG-ADL also showed no significant difference 12 weeks (T3) after vaccination in the MG vaccination group compared to T0 and T1 ($p=0.12$). In the placebo group there was no significant difference between any of the 4 time points at which the MG-ADL was performed (T0-T3) (data not shown).

3.5 Antibodies against AChR

No change in antibody titre was observed 4 weeks after influenza vaccination (Figure 3).

3.6 Side effects

The MG vaccination group reported side effects in 30.4% (7/23) at T1, the placebo group in 37.5% (9/24) at T1 ($p=0.6$). At T2, 4 weeks after unblinded influenza vaccination of the placebo group 52% (12/23) reported side effects. At T1 healthy controls reported significantly more side effects (70%; 33/42) than the MG vaccination or placebo group ($p<0.01$). The most commonly reported side effects for MG or HC were local redness and soreness at the injection site. No change in MG symptoms was reported in the MG group at T1. In the placebo group, 3 patients reported a mild exacerbation of their MG symptoms during the T1-T2 period. Exacerbation of symptoms lasted 1 day to 1 week after vaccination and did not lead to a change in medication.

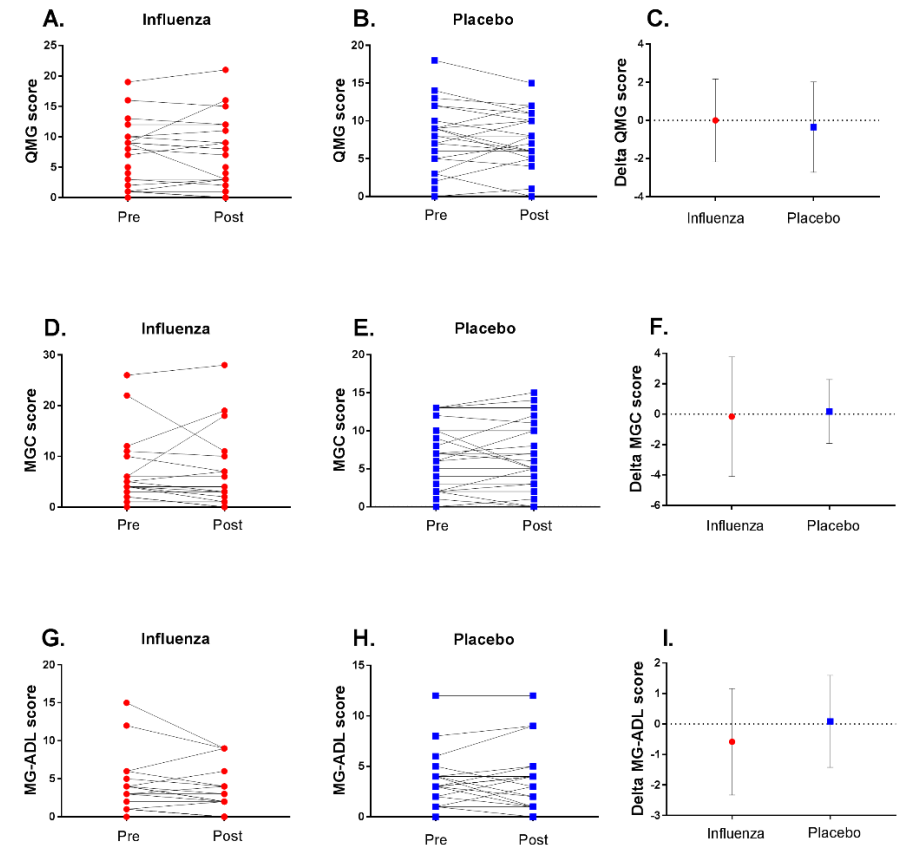


Figure 4. Clinical outcome measures for the AChR MG vaccination group (●) and placebo group (■) pre- and 4 weeks post-vaccination of the Quantitative Myasthenia Gravis score (QMG), Myasthenia Gravis Composite score (MGC) and Myasthenia Gravis Activities of Daily Living (MG-ADL) in A, B, D, E, G and H. The delta of the clinical outcome scores are shown in C, F and I. No significant differences were found.

4. DISCUSSION

In this prospective, double blind, randomized, placebo-controlled study we show that in AChR MG patients influenza vaccination is safe and induces an immune response comparable to that of healthy controls. The study population consisted of patients with stable disease and a stable medication regime in the past 3 months. A seroprotective titre for all three strains of the seasonal influenza vaccine was reached in 40.4% (19/47) of the AChR MG group and in 51% (24/47) of the HC group. IM or thymectomy status did not significantly influence post vaccination GMT titres. No clinical or immunological exacerbation was found as

clinical outcome scores and AChR antibody titres showed no significant changes. It is generally assumed that patients with an autoimmune disease are more prone to infections, resulting in increased morbidity and mortality [2]. In autoimmune inflammatory rheumatic disease, influenza-vaccinated patients have a lower incidence of pneumonitis, acute bronchitis and viral infections than unvaccinated patients [3]. To our knowledge no such studies have been performed in patients with MG. Recently a randomized controlled trial on influenza vaccination showed that influenza vaccination is safe, based on QMG scores and AChR antibody titres, but without including an healthy control group [8]. Studies on the efficacy of influenza vaccination in rheumatic disease also found that achievement of seroprotection (post HI-titre $\geq 1:40$) is similar to healthy controls, irrespective of medication [3]. In patients with SLE, the response to influenza vaccination is comparable to that of healthy controls [3]. Two studies showed a trend towards a lower response to vaccination in patients who used azathioprine [20, 21], which is also commonly used in MG next to corticosteroids. In this study we did not find a significant effect of IM on the humoral response. Due to small size of treatment subgroups and because of frequent combinations of IM, we could not investigate specific effects of a single drug. In a study on the efficacy and safety of a tetanus vaccination in MG, we found that IM lowers pre- and post-vaccination GMTs, but did not affect the efficacy of the response [9]. This difference might be explained by the type of vaccine that is investigated and the vaccination history of the patients.

Some MG patients chose not to participate out of concern for an exacerbation of their symptoms. Even in our trial participants, only two-thirds had obtained an influenza vaccination in previous years, similar to the frequency of our healthy controls. The tetanus revaccination study in AChR MG patients showed a small but statistically significant increase of the QMG score of 1 point at 4 weeks, which is far less than the 2.3 points that is generally accepted as the minimal clinically relevant difference. A recent study indicated that an exacerbation of MG is more likely after an influenza-like infection or a common cold, than following an influenza vaccination (10/25 (40%) and 15/96 (15.6%) vs. 2/133 (1.5%) [7]. In line with our results, no clinical exacerbation was found in patients with RA and SLE following influenza vaccination [3]. Interestingly, unblinded influenza vaccination of MG patients in T1-T2 resulted in more reported side effects and a higher incidence of self-reported aggravation of MG symptoms than blinded vaccination or placebo injection. This may be explained by the presence of a prejudice among MG patients that vaccination might be harmful, leading to increased reporting of subjective complaints.

4.1 Strengths and limitations

The main strengths of this study are its placebo-controlled, double blind, randomized design and the systematic assessment of multiple relevant measures of clinical disease severity at multiple time points up to twelve weeks.

Limitations are the exclusion of patients with severe or unstable MG and patients using high doses of corticosteroids. Therefore, we cannot draw a conclusion on

the safety and efficacy of vaccination in these groups. Although the study was not powered to detect small changes in clinical outcomes, none of these measures show a trend indicating a possible negative effect.

Theoretically, the unblinded nurses may have caused unblinding of patients, but they specifically ensured that patient blinding was maintained during injection. Furthermore, clinical outcome measures, which are likely the most susceptible to unblinding were taken before unblinding the patients 4 weeks after the injection. Median age of healthy controls was lower, which might result in a stronger humoral response. However, no significant post-vaccination differences were observed between MG and HC groups.

5. CONCLUSION

The antibody response to an influenza vaccination in patients with mild to moderate MG is similar as in healthy subjects, and not affected by the use of immunosuppressive medication. Influenza vaccination did not induce any immunological or clinical exacerbation of MG.

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