Phylogenetic analysis of the neuraminidase segment gene of Influenza A/H1N1 strains isolated from Monastir Region (Tunisia) during the 2017–2018 outbreak

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Abstract
Influenza A/H1N1 is widely considered to be a very evolutionary virus causing major public health problems. Since the pandemic of 2009, there has been a rapid rise in human Influenza virus characterization. However, little data is available in Tunisia regarding its genetic evolution. In light of this fact, our paper aim is to genetically characterize the Neuraminidase, known as the target of antiviral inhibitors, in Tunisian isolates circulating in Monastir region during 2017–2018. In total of 31 positive Influenza A/H1N1 detected by multiplex real-time PCR, RT-PCR of neuraminidase was performed. Among the 31 positive samples, 7 samples representing fatal and most severe cases were conducted for sequencing and genetic analysis. The results thus obtained showed genetic evolution of the A/H1N1 neuraminidase between 2009 and 2010 and 2018–2019 outbreaks. All Tunisian isolates were genetically related to the recommended vaccine strain with a specific evolution. Moreover, the phylogenetic analysis demonstrated that France and especially Italian strains were the major related strains. Interestingly, our results revealed a specific cluster of Tunisian isolates where two intragroup were evolved in correlation with the severity and the fatalities cases. From the outcome of our investigation, this study confirms the genetic evolution of the Influenza A virus circulating in Tunisia and gives a preliminary analysis for a better comprehension of new emerging Tunisian strain’s virulence and thus, a more appropriate monitoring of Influenza virus A/H1N1 during each round of outbreaks.

Keywords Influenza A/H1N1 · Neuraminidase · Phylogenetic analysis · Severe acute respiratory infections

Introduction
Seasonal Influenza vaccines are the key public health tool against the Influenza virus. However, its efficacy remains limited due to the antigen mismatch between vaccine strains and circulating viruses. The immune status of the host can lower the effectiveness which varies by age group and by the vaccination status (Osterholm et al. 2012; Lewnard and Cobey 2018). Seasonal Influenza epidemics cause up to 650,000 respiratory deaths and 3 to 5 million cases of severe illness per year worldwide and present a real economical problem (Petrova and Russell 2018). Influenza viruses are segmented single-stranded, negative-sense RNA viruses that belong to the Orthomyxoviridae family. Based on their antigenic specificity of the nucleoprotein and matrix protein, Influenza viruses are divided into Influenza A (IAV), Influenza B,
and Influenza C (Bouvier and Palese 2008). Several publications have appeared in recent years documenting that the IAV virus exhibits the most circulating type causing disease in humans, and different animal and avian species (Horimoto and Kawaoka 2005). The Influenza genome contains 8 RNA segments (PB2, PB1, PA, HA, NP, NA, M, and NS). The 6th segment encodes for the Neuraminidase (NA), one of the major surface glycoproteins of the Influenza virus (Bouvier and Palese 2008). There are 11 different neuraminidase subtypes found in circulating IAV (N1 through N11, respectively). Only N1 and N2 which are expressed in H1N1 and H3N2 subtypes cause endemic in humans and give rise to seasonal outbreaks (Mcauley et al. 2019). In contrast to IAV which is classified into subtypes, Influenza B is classified into two lineages: B/Yamagata and B/Victoria (Hatta and Kawaoka 2003). The IAV can be further classified into specific clades and subclades. For example, IAV (H1N1) belongs to the clade 6B1 and the subtype 6B1.A (CDC 2019). It has now been demonstrated that NA plays several roles during the infection process. It acts in the early stage of infection by enhancing virus entry contributing to an efficient virus-cell association (Guo et al. 2018). In addition, this glycoprotein acts in the final stage of infection by removing sialic acid from cellular glycoconjugates in a specific site. Thus, newly synthesized virions are released from the infected cell to spread the infection (Palese et al. 1974). During the virus infection, and to escape from the immune system, the natural selection favors the continuous modifications of the antigenic structure of NA by generating mutations. Since NA is an important antiviral target, a single substitution on a specific site can lead to a resistance capacity against NA inhibitors such as zanamivir and oseltamivir (Hurt et al. 2012; Pinilla et al. 2012; Krammer et al. 2018).

In the last few years, there has been a growing interest in the genetic variability of Influenza NA to understand the causes of the immune escape and its impact on the fitness. However, in Tunisia, studies on genetic characterization of Influenza NA are still lacking. In this paper, we focus on Influenza strains. For example, IAV (H1N1) belongs to the clade 6B1 and the subtype 6B1.A (CDC 2019). It has now been demonstrated that NA plays several roles during the infection process. It acts in the early stage of infection by enhancing virus entry contributing to an efficient virus-cell association (Guo et al. 2018). In addition, this glycoprotein acts in the final stage of infection by removing sialic acid from cellular glycoconjugates in a specific site. Thus, newly synthesized virions are released from the infected cell to spread the infection (Palese et al. 1974). During the virus infection, and to escape from the immune system, the natural selection favors the continuous modifications of the antigenic structure of NA by generating mutations. Since NA is an important antiviral target, a single substitution on a specific site can lead to a resistance capacity against NA inhibitors such as zanamivir and oseltamivir (Hurt et al. 2012; Pinilla et al. 2012; Krammer et al. 2018).

In this context, the present work aims to study the virus phylogeny and to explore the evolutionary processes of NA that shape the genetic diversity of Tunisian IAV(H1N1) strains.

**Materials and methods**

**Ethics approval**

Samples were collected after the receiving of written informed consent from the subjects or their legal representatives. The study protocol was approved by the research ethics committee of the faculty of medicine, the university of Monastir under the number [IORG0009738 N° 18/ OMB0990-0279].

**Samples and clinic data collection**

During the 2017–2018 Influenza outbreak, nasopharyngeal swab samples were collected from patients suffering from severe acute respiratory illness in the Sahel region of Tunisia (Monastir) in collaboration with the laboratory of medical microbiology of Fatouma Bourguiba university hospital Monastir, Tunisia. In this work, we focus on 31 samples that were detected positive for A/H1N1 by multiplex real-time PCR (RespiFinder® 2smart PathoFinder) as per the manufacturer’s instructions, during respiratory virus routine detection. Each sample was received with its clinic information then stored directly at -80°C for RT-PCR based detection of NA and genetic analysis. All infected patients were not vaccinated against the Influenza virus.

**Viral RNA extraction and RT-PCR of NA gene**

RNA extraction from clinical samples was performed using TRIzol™ Reagent (Invitrogen™) according to the manufacturer’s instructions.

The cDNA synthesis of the whole Influenza genome was performed using the universal primer UNIT12 (AGCRAAAGCAGG) (Hoffmann et al. 2001), the viral RNA was mixed with 20 pMol of Unit 12 primer and 12.5 pM of dNTP for a pre-heating step at 65°C during 5 min. The mixture was incubated immediately on ice. 10 U/µl of Mmlv enzyme (Biogene) and 10 µM of DTT (dithiothreitol) was added to the previous mixture and the total volume was incubated for 60 min at 42°C for the retro-transcription.

PCR reaction for the neuraminidase gene amplification was performed by TaKaRa LA Taq® DNA-Polymerase using specific primers (NA F:5’AGCAGGAGTTAAAATGAAT CCAAA3’) and (NA-R:5’CTTCCTATCCAAAACACCAC T3’) for NA of seasonal IAV (H1N1) amplification (1097pb) (WHO 2017a).
The thermal cycle was programmed for amplification as follows: Incubation at 95°C for 1 min, and then 30 cycles of denaturation at 94°C for 30', 58°C for 30' for hybridization, 72°C for 2 min followed by a final elongation step at 72°C for 5 min using a thermocycler 2770 (Applied Biosystems).

An agarose gel (1 %) in 1X Tris-borate-EDTA and ethidium bromide (250 ng/μl) was prepared. Then, 5 μL of NA PCR amplification products were added to 1 μL of 6X loading and loaded. 3 μL of 1 kb DNA ladder (GeneON) were also loaded. Finally, the gel was run at 100 Volts for 30 then NA bands were visualized and photographed under UV light using « Quantity One » software of Gel Doc 2000 (Bio-Rad).

**PCR products purification and sequencing**

Since we aim to sequence the NA segment and analyze it genetically, the key step is to have a good quality of DNA from the RT-PCR for the NA. Thus, we kept only samples where nasopharyngeal swabs were collected during the first 5 days as the recommendation of the centers for disease control and prevention (CDC 2020). Also, samples from fatal cases were conducted for sequencing to study the virulence of their strains (Methodology is shown in Fig. 1).

The 7 selected PCR amplification products were purified from PCR reagents following the clean-up protocol steps from FavorPrep GEL/ PCR purification Kit as per the manufacturer’s instructions. These 7 samples were subjected for sequencing (commercially) using the big dye terminator method in 3500 ABI Prism Genetic Analyzer (Applied Biosystems) with the same NA primers recommended by the WHO for NA detection and sequencing studies.

**Nucleotide sequence submission**

The nucleotide sequences data obtained from this study have been submitted to NCBI GenBank. The accession number of the sequences for the NA gene obtained are MT239356 to MT239362.

**Phylogenetic analysis**

Tunisian sequences were aligned using Cluster X software (Larkin et al. 2007) with other European, and Mediterranean strains sequences obtained randomly from online Influenza database from the 2009–2010 season until the 2018–2019 season available on [https://www.ncbi.nlm.nih.gov/genomes/FLU/Database/nph-select.cgi?go=genomeset]. WHO recommended vaccine strain A/Michigan/45/2015 (A/H1N1) pdm09-like for the 2017–2018 season was also included in the alignment. The phylogenetic tree was generated in MEGA version 7.0 (Kumar et al. 2016) using the maximum likelihood method with the general time-reversible (GTR) + G evolutionary model. 1000 bootstrap replicates were used to construct the phylogenetic tree and high bootstrap values above 70 % were visualized. The final tree was visualized and annotated with FigTree v.1.4.2 (http://tree.bio.ed.ac.uk/software/figtree).

![Overview of the methodology used to characterize Neuraminidase gene of Influenza A (H1N1) in Monastir region (Tunisia)](image-url)}
Results

Clinical characteristics

Demographic data

The study included data generated from 31 samples in the region of Monastir, Tunisia. The population was composed of 31.3% males and 38.7% females. The patient’s ages ranged from 3 months to 80 years with a mean age of 38.16 ± 31.46. The age group with the highest number of SARI was the 0–2 years with (36.5%), followed by the ≥65 years with 29%, lastly the 36–64 years and 20–35 with a rate of 19.4%, 12.9% respectively. Among the 31 positive cases, 19 patients were hospitalized in ICU (61.3%) and 12 patients (38%). 3 of the 31 patients are passed away (9.68%) which the ages are ranged from 3 months to 2 years (Table 1). Taking into consideration the fatality cases, the most severe cases, and the period between the first sign of infection (days ≤5 days), only 7 positive samples were sequenced which represents (19.35%).

Clinical outcomes in the totality of infected patients

The clinical outcomes of IAV (H1N1) are shown in Table 2. The majority of patients suffered from fever (96.77%) and cough (24%). In total, 22.58% of patients developed respiratory distress and dyspnea 32.25%. In total 10/31 (32.25%) of patients received antiviral treatment and 9.67% of patients received mechanical ventilation. In addition to the respiratory outcomes, the infection was associated with comorbidity (22.58%) like diabetes, hypertension, cardiovascular diseases. From the 31 patients, 22.58% have immunodeficiency and 6.45% had a pregnancy."

Clinical data of the selected sequenced samples

The genetically analyzed infected cases were predominantly males (6 male and 1 female) aged between 3 months to 70 years. All samples were taken from patients suffering from SARI coming from the Monastir region. Sampling was performed in the 2017/2018 season (One sample during the first trimester and 6 during the second trimester). All the analyzed 7 patients presented Influenza A symptoms (caugh, fever, fatigue), Besides, the 3 fatal cases (SB. G51; SB. G47; SB. G67) were suffering from additional respiratory complications (distress). Real-time RT-PCR revealed IAV (H1N1) as a responsible agent of these acute respiratory infections for all the samples with one case of co-infections with adenovirus. Only 2 patients SB. G70 and SB.G47 were treated with oseltamivir. The overall clinical characteristics of the studied samples are summarized in Table 3.

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**Table 1** Demographic data of the positive Influenza A (H1N1) samples

| Region                        | Monastir, Tunisia | 31 (100%) |
|-------------------------------|-------------------|-----------|
| Gender                        | Males             | 19 (61.3%)|
|                               | Females           | 12(38.7%) |
| Age (years)*                  | Mean              | 38.16±31.46|
|                               | [0–2]             | 11(35.5%) |
|                               | [20–35]           | 4(12.9%)  |
|                               | [36–64]           | 6(19.4%)  |
|                               | ≥65               | 9(29%)    |
| ICU admission                 | Yes               | 19 (61.3%)|
|                               | No                | 12(38%)   |
| Post infection state          | Fatality          | 3 (9.68%) |
|                               | Recovery          | 27 (90.32) |
| Days between first sign of infection and sampling | ≤5 days             | 19,35%   |
|                               | >5days            | 80.65%    |

*No patient belong to the age group [3–19]
Sequencing

Sequences obtained for all amplifications were verified, aligned, and completed manually using reference sequences from the GenBank. The accession numbers of the sequences for the NA gene obtained were registered in the GenBank with accession numbers from MT239356 to MT239362.

Phylogenetic analysis

As can be seen from Fig. 2, phylogenetic analysis for the NA gene of IAV(H1N1) Tunisian isolates highlighted two major clades: Clade A and Clade B. On one hand, the NA sequences analysis revealed that all Tunisian strains of the 2017–2018 season had evolved away from IAV(H1N1) pandemic strains circulating during the 2009–2010 and 2010–2011 outbreak (clade A). In contrast, they belonged to clade B with the vaccine strain A/Michigan/45/2015 (A/H1N1) pdm09-like and European IAV strains circulating during the 2017–2018 season in several Mediterranean countries: Spain, Italy, and France. On the other hand, the phylogenetic tree showed clearly that IAV(H1N1) Tunisian isolates were related to the vaccine strain A/Michigan/45/2015 (A/H1N1) pdm09-like virus recommended by the WHO for the northern hemisphere during 2017–2018. However, our isolates showed a genetic variation when related to this vaccine strain.

Besides, it has been found that Tunisian isolates clustered with IAV strains circulating during the 2018–2019 outbreak such as the Italian strain IAV (A/Italy/7841/2019 H1N1, A/France/180130-4/2018 H1N1, and A/France/180130-2/2018 H1N1) forming together a distinct clade: clade B1 belonging to the clade B (Fig. 2) where they share the same ancestor.

Focusing on Tunisian isolates strains themselves, our results revealed two different intragroup clades B.1.1 and clade B.1.2 in the same population hospitalized in Fatouma Bourguiba Monastir for SARI infections during Influenza 2017–2018 outbreak. Interestingly, the phylogenetic tree demonstrated a proper cluster where mortality cases were assembled. However, other severe cases were assembled in another cluster. These results offer a powerful correlation between phenotypes and genotypes. As far as we know, this is the first time that the NA of Tunisian IAV(H1N1) isolates causing fatalities in the Sahel region were characterized.

Discussion

In this study, we focused on severe acute respiratory illness caused by IAV(H1N1) to investigate the reason for fatal and severe infections. Usually, IAV(H1N1) was described to be associated with fever and coughs symptoms. When infecting
IAV(H1N1) can cause inflammation of the upper and lower respiratory tract (Taubenberger and Morens 2008; Jilani et al. 2020). The infection period persists for three to seven days for healthy persons. However, it can be longer (9 to 10 days) for patients with high risk such as children, pregnant women, and aged subjects (Reed et al. 2014). Thus, the inflammation will be associated with further complications such as respiratory distress and dyspnea requiring in most cases intensive care and antiviral treatments. The highest infected age group in all patients was children ≤ 2 years with 3 fatal cases which explains the high risk of infection for this age category. This description supports a previous Tunisian study showing the severity of IAV(H1N1) infection outcomes in hospitalized children with a median age of 12 months (Tinsa et al. 2018). The most likely explanation about infection susceptibility of young children is their fragile immune system which has never been exposed to IAV previously (Pinilla et al. 2012). So, their organisms can develop more severe symptoms and shed a larger number of viruses for a longer time. Besides, the second-highest age group developing SARI infection is the ≥ 65 years confirming the great prevalence of this category of age to be infected and hospitalized for IAV(H1N1) (Taubenberger and Morens 2008).

In this paper, our attention was directed to only 31 SARI caused by IAV(H1N1) in the Monastir region. These patients were enrolled in the university hospital of Fatouma Bourguiba in several departments (polyvalent ICU, pediatrics department, emergency department, infection diseases department, and maternity ICU). However, non-Influenza respiratory viruses can be also a serious emerging threat of respiratory infection during the same season. In Tunisia (Sousse Area), it has been demonstrated that the rhinoviruses, respiratory syncytial virus A/B, adenovirus, human coronavirus (NL63, HKU1, OC43, and 229) are the most frequently detected viruses in children. In addition, IAV (H3N2) and can circulate in the same period and cause similar infections (Brini et al. 2017).

The time of sampling is critical for IAV diagnostics. After 5 days of symptoms appearing, the virus titer decreases in the upper respiratory tract which makes the qualitative PCR detection of NA from nasopharyngeal swabs complicated. Thus, only a few samples representing (19.35 %) from the totality were sequenced and analyzed genetically (3 fatal cases, 4 most severe cases).

From the studied samples, we reported fatal cases. We suppose that, once the virus binds to columnar epithelial cells in the respiratory tract, the virus had a high ability of replication leading to a high virus titer. Thus, the Influenza virus may cause severe diseases including apoptosis of the host and adjacent cells origin of respiratory complications described in Table 3 (Otte et al. 2016). From the two treated subjects with NA inhibitor (oseltamivir), one died (SB.47). For that reason, we focused on NA amplification and sequencing for more genetic exploration. Interestingly, the amplification of the NA gene of seasonal IAV(H1N1) has been poorly realized in Tunisia and most interest has been focused on the Hemaglutunine gene for detection and characterization (El Moussi et al. 2013; Ben M'hadheb et al. 2015; Soli et al. 2019).

Although all the patients included in this study were detected positive for IAV (H1N1) by real-time PCR, the qualitative RT-PCR for the neuraminidase showed some weak bands. This can be explained by the time from illness onset to specimen collection, and the type of the sample. To maximize detection of Influenza viruses, respiratory specimens should be collected as close to illness onset as possible (ideally < 3–4 days). However, among the 31 samples only (19.35 %) were sampled ideally. Molecular assays may detect Influenza viral RNA in respiratory tract specimens for longer periods after illness but re-using the same sample for qualitative RT-PCR might impact the quality of bands, and therefore, sequencing.

The phylogenetic study demonstrated a divergence between the pandemic IAV(H1N1) strains isolated in 2009 and 2010 and the Tunisian isolates during 2017–2018 forming a defined clade. The main clades were organized regarding the pandemic strain to a seasonal Influenza virus. It’s the antigenic drift phenomenon (Su et al. 2015; Petrova and Russell 2018).

Again, our data shows clearly that all the Tunisian isolates of IAV(H1N1) included in this study, share the same ancestor with the vaccine strain. Every year, the WHO design an Influenza vaccine for the Northern and Southern Hemisphere. During the 2017–2018 season, the vaccine composition included A/Michigan/45/2015 (A/H1N1) pdm09-like virus (WHO 2017). In the present study, we focus on the IAV(H1N1) subtype because of its predominance and severity during this season in North Africa and specifically in Tunisia.
As mentioned above, our results concur well with the genetic characterization of 32 European IAV(H1N1) strains during 2017–2018 which showed a grouping to the sub-clade 6B1 presented by the vaccine A/Michigan/45/2015 (WHO 2018b). Besides, it has been reported, globally, that IAV(H1N1) viruses are in genetic flux, with substantial heterogeneity declaring a fitness advantage (ECDC 2018). In line with this, the genetic diversity in Europe did not seem to affect the vaccine effectiveness against IAV(H1N1)pdm09 in most studies (Rondy et al. 2018). Interestingly, the analysis for the NA characterization of Tunisian IAV(H1N1) isolates has revealed genetic variations compared to the vaccine strain although they are genetically related. Thus, a probable NA evolution has occurred.

Clade B1 has gathered, mostly, Mediterranean IAV(H1N1) strains circulating in the same period 2017–2018. Tunisian isolates were closely related especially to French and Italian strains. This genetic relation seems to be explained by the virus transmission between individuals in Europe and individuals in Tunisia due to the geographic position. Since Tunisia is located in a central position between North Africa and Europe, frequent movement of individuals, especially between France and Italy, might promote virus transmission. Tourism activity, close economic relationships can argue this hypothesis (France diplomacy 2017). During 2018 the national Tunisian office of tourism reported that 30,8 % of tourists are coming from Europe (ONTT 2018).

Our attention was focused not only on the comparison of the Tunisian isolates with the Mediterranean strains but also on the phylogeny between the circulating viruses in the same Tunisian population. Particular attention is paid to the shape of the tree describing two internal clusters. The remarkable result to emerge from this data is that internal clusters have been distinguished regarding the severe infection effect: fatal cases together, and other severe cases together. Thus, IAV(H1N1) circulating in the Monastir region of Tunisia have different virulence and severity between patients. Our study provides additional support for a possible correlation between the phenotype and genotype that has never been described in most studies (Rondy et al. 2018). Interestingly, the analysis for the NA characterization of Tunisian IAV(H1N1) isolates has revealed genetic variations compared to the vaccine strain although they are genetically related. Thus, a probable NA evolution has occurred.

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The main limitation of this study is the lack of NA sequences in the Influenza database in some Mediterranean regions during 2014–2015 and 2015–2016. Also, a deeper investigation of possible mutations and aminoacid alterations would be very interesting if the sequence covers fully the neuraminidase segment. Moreover, the few numbers of Influenza virus Tunisian isolates seem to present a further limitation of this study.

This work is the first step of IAV(H1N1) genetic evolution understanding in the region of Monastir, Tunisia. The NA characterization will certainly help to solve virus resistance phenomena during antiviral treatment, could potentially help on the monitoring of Influenza virus circulation in Tunisia, and might estimate vaccine efficiency against new emerging strains.

Conclusions

From the research that has been carried out, it is possible to conclude that Tunisian IAV(H1N1) isolated in the Monastir region are genetically related to France strains and especially to Italian strains. A possible genetic heterogeneity and recombination events between the Tunisian circulating strains 2017–2018 might be the origin of such virulent strains. Our results are promising and should be continued with whole-genome sequencing to study deeply the process of evolution in terms of mutations on the genetic and proteomic levels.

Abbreviations. IAV, Influenza A virus; NA, Neuraminidase; SARI, Severe acute respiratory infections; NCBI, National Center for Biotechnology Information; WHO, World Health Organization; PCR, Polymerase chain reaction; HA, Hemagglutinin

Supplementary Information. The online version contains supplementary material available at https://doi.org/10.1007/s11756-021-00723-y.

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Author contributions. Conceptualization: [Sabrine Ben Hamed, Jawhar Gharbi]; Methodology: [Sabrine Ben Hamed, Myriam Harrabi]. Formal analysis and investigation: [Sabrine Ben Hamed, Aida Elargoub,
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**Declarations**

**Ethical standards** Human Nasopharyngeal swabs samples used in this study were obtained by special physicians for the routine laboratory testing of severe acute respiratory illness using real-time RT-PCR. The study protocol was approved with a formal authorization by the Research Ethics Committee, Faculty of Medicine of Monastir, under the number [IORG0009738 N° 18/OMB0990-0279].

**Conflict of interest** The authors declare no conflict of interest.

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