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ORIGINAL ARTICLE

Adherence to Anaphylaxis Guidelines: Real-World Data From the Emergency Department of a Tertiary Hospital

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Abstract

Background: Few studies have evaluated adherence to anaphylaxis guidelines in emergency departments (EDs). **Objective:** The objective of this study was to evaluate adherence to anaphylaxis guidelines in the ED of a tertiary hospital. **Methods:** Medical records of patients attended in the ED of University Hospital of Salamanca, Spain were reviewed. Those patients fulfilling the anaphylaxis criteria proposed by the NIAID/FAAN were selected. **Results:** A total of 89 patients (47 adults and 15 children), all with anaphylaxis reaction, were included. The anaphylactic reaction was moderate in 65% of adults, severe in 34%, and very severe in 1%. In children, all reactions were moderate. Fewer than half of the patients (42%) received adrenaline in the ED; this was administered intramuscularly in only 19% of cases. As for the severity of the reaction, 65% of patients with moderate reactions and 42% with severe reactions were not treated with adrenaline. At discharge from the ED, an auto-injector was prescribed in 6.6% of adults and 1.6% of children. Fifty-four percent of patients attended alone (57% adults vs 27% children, $P=0.047$), 29% instructions about avoidance of triggers (31% adults vs 20% children, NS), and 51% written instructions for recognition of anaphylaxis were provided. **Conclusion:** The results of this study reveal a significant discrepancy between recommendations in guidelines and management of anaphylaxis in the ED. Additional training efforts are needed to improve the treatment of patients with anaphylactic reactions.

Key words: Anaphylaxis, Guidelines, Adrenaline.

Resumen

Antecedentes: Pocos estudios han evaluado el cumplimiento de las recomendaciones de las guías clínicas de anafilaxia en los servicios de urgencias. **Objetivo:** El objetivo de este estudio fue conocer el cumplimiento de las guías de anafilaxia en el servicio de urgencias (SU) de un hospital terciario. **Métodos:** Se revisaron los informes de los pacientes atendidos en el SU del Hospital Universitario de Salamanca durante un año y se seleccionaron los que cumplían los criterios de anafilaxia propuestos por el NIAID/FAAN. **Resultados:** Se identificaron 89 pacientes (47 adultos y 15 niños). El 65% de los adultos presentó una reacción moderada, el 34% grave y el 1% muy grave; en todos los niños la reacción fue moderada. Menos de la mitad de los pacientes (42%) fueron tratados con adrenalina en el ED; esta se administró intramuscularmente en solo 19% de los casos. En lo que respecta a la severidad de la reacción, el 65% de los pacientes con reacciones moderadas y el 42% con reacciones severas no recibieron adrenalina. Al alta, se recomendó un auto-inyector de adrenalina al 6.6% de los pacientes, se dieron indicaciones para evitar posibles desencadenantes al 29% (31% adultos frente a 20% niños, $P=0.047$) y se proporcionaron instrucciones escritas para reconocer los signos de alarma de una reacción anafiláctica al 51% (41% adultos frente a 100% niños, $P=0.001$). **Conclusión:** Los resultados de este estudio revelan una discrepancia significativa entre las recomendaciones en las guías clínicas y el manejo de la anafilaxia en el ED. Es necesario un mayor esfuerzo en la educación para mejorar el tratamiento de los pacientes con anafilaxia.

Palabras clave: Anafilaxia, Guías clínicas, Adrenalina.

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**ACTION PLAN FOR
Anaphylaxis**
 For use with adrenaline (epinephrine) autoinjectors

Name: _____ Date of birth: _____

Confirmed allergies: _____

Family/emergency contact name: _____

Work Pno: _____ Home Pno: _____ Mobile Pno: _____

Plan prepared by medical or nurse practitioner: _____

I hereby authorise healthcare providers specified on this plan to administer this plan to me.

Signed: _____ Date: _____ Action Plan due for review - date: _____

How to give EpiPen® adrenaline (epinephrine) autoinjectors

1 Turn hot around Eddies and PULL OFF BLUE SAFETY RELEASE

2 Hold with one PLACE ORANGE END against outer thigh (left or right) and push firmly

3 PUSH DOWN HARDS until a click is heard and hold in place for 3 seconds REMOVE EpiPen

EpiPen® is prescribed for children over 20kg and adults. EpiPen® is prescribed for children 20-2kg.

If EpiPen® is prescribed for children under 20kg, please follow the instructions on the prescription.

© ASCIA 2018 This plan may be copied if it is completed and signed by the patient's medical or nurse practitioner and cannot be altered without their permission.

Emergency treatment for a suspected anaphylactic reaction

Investigation in adults or young people aged 16 years or older

Take timed blood samples for mast cell tryptase testing:
 as soon as possible after emergency treatment
 ideally within 1–2 hours (but no later than 4 hours) from the onset of symptoms.
 Inform the person (or, as appropriate, their parent and/or carer) that a blood sample may be required at follow-up with the specialist allergy service to measure baseline mast cell tryptase.

Investigation in children younger than 16 years

Consider taking blood samples for mast cell tryptase testing if the cause is thought to be venom-related, drug-related or idiopathic:
 as soon as possible after emergency treatment
 ideally within 1–2 hours (but no later than 4 hours) from the onset of symptoms.
 Inform the parent and/or carer that a blood sample may be required at follow-up with the specialist allergy service to measure baseline mast cell tryptase.

Assessment

Document the acute clinical features of the reaction:
 rapidly developing, life-threatening problems involving the airway (pharyngeal or laryngeal oedema), and/or
 hæmorrhage (haemorrhage with tachycardia) and/or

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