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Adrenaline Autoinjector Needle Interlocking in the Thumb Due to Improper Injection

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Abstract: Herein, a case of improper epinephrine autoinjector injection into the thumb involving deep connective tissue and bone is reported. We additionally review knowledge, skills and habits of a cohort of autoinjector users of a hospital-based dermatologic outpatient clinic.

Keywords: Epinephrine autoinjector, digital ischemia, application error.

INTRODUCTION

Accidental digital injection of epinephrine autoinjectors is a known side-effect when dispensing this units to patients suffering from anaphylactic reactions [1]. In this case story, we report a 42-year old man with hymenoptera venom allergy and a history of a systemic reaction. The patient had been evaluated in an outpatient allergy clinic. According to current guidelines [2], he had been equipped with an emergency medication, including an epinephrine autoinjector (EpiPen®) for self-treatment of allergic reactions. He had been instructed in the proper use when he was issued the epinephrine autoinjector (EpiPen[®]) by a dermatologist, as according to a long-standing tradition, in Austria, patients with hymenoptera venom allergy are evaluated and treated by dermatologists.

After being stung by a yellow jacket one day, the patient experienced a sensation of impending collapse. He attempted to employ the autoinjector, but, after unlocking he gripped it upside down causing the needle to penetrate his right thumb's distal phalanx. Within minutes, the patient was brought to the emergency department by the emergency physician who had already administered intravenous antihistamines and corticosteroids. On arrival, the patient had already recovered from the symptoms of an allergic reaction, but his thumb was found to be pale and cold with the autoinjector still stuck in the patient's thumb. An X-ray of the hand revealed the needle being located in the soft tissue, furthermore the tip of the needle had hit the bone and had been bent more than 90 degrees (Fig. 1a). Not only the mechanism had deployed as expected and injected the epinephrine dose into the thumb, but the needle stuck in the thumb acting as a barbed hook (Fig. 1b). By careful manoeuvering, controlled by sequential radiograms, the needle was removed. Subsequently, the patient received topical application of glyceryl trinitrate and concomitant oral administration of acetylsalicylic acid. Local infiltration of the β_2 agonist terbutaline, which had been suggested by the Vienna poison center, was refused by the patient. Finally, the impaired circulation could be restored. There was no permanent structural or functional damage observed in the further clinical course.

With our interest drawn to incorrect epinephrine autoinjector use, we made a short interview with a cohort of 19 patients diagnosed with hymenoptera venom allergy, attending our clinic for desensitization therapy with hymenoptera venom (Table 1). Over a period of three months, 17 of 19 interviewed patients were still carrying the autoinjector (Epipen®) with them (89.5%). In three of the remaining 17 cases (17.6%) the autoinjector had expired, leaving 14 patients equipped with a well functioning autoinjector (82.35%). We investigated whether the patients were still able to properly use a demonstrator autoinjector pen. Ten of 14 patients with a well functioning autoinjector (71.4%) were capable to use the demonstration pen in a correct manner; in 4 cases the use was incorrect including one ectopic application, one application of too short contact duration (no injection triggered) and two incorrect applications for other reasons. The three patients with an expired autoinjector were not capable to demonstrate the autoinjector use in a correct way, including one ectopic application, one application of too short contact duration (no injection triggered), and one incorrect application of other reason (cap not removed and therefore no injection triggered).

Several cases of accidental injection of epinephrine autoinjectors into digits including the thumb have been published. The estimated incidence is about one accidental digital injection per 50 000 autoinjector units [3]. Epinephrine causes vasoconstriction, the harmed digit is at risk for necrosis. Suggested treatment includes immersion in warm water, topical nitroglycerin, local injection of terbutaline [4]

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Fig. (1a). X-ray of the epinephrine autoinjector locked in the distal phalanx of the right thumb.



Fig. (1b). The distorted needle of the epinephrine autoinjector (Epipen®, ALK-Abelló, Linz, Austria) after removal from the thumb.

Table 1.	Patient Characteristics and Test Results. Abb	previations: F: Functional	l Autoinjector, E: Expired	l Autoinjector, C: Correct
	Application, I: Incorrect Application, N: Autoi	injector not Available		

Patient	Age	Sex	Allergy	Classification of Allergic Reactions [11]	Use of Training Device
1	45	m	Yellow jacket	IV	F,C
2	38	m	Yellow jacket	II	F,C
3	67	f	Yellow jacket	IV	F,C
4	60	m	Yellow jacket	IV	F,C
5	56	f	Yellow jacket	IV	F,I
6	67	f	Honeybee, Yellow jacket	IV	E,I
7	62	f	Yellow jacket	Ш	F,C

Patient	Age	Sex	Allergy	Classification of Allergic Reactions [11]	Use of Training Device
8	54	f	Yellow jacket	IV	F,I
9	64	f	Yellow jacket	III	F,I
10	63	f	Yellow jacket	П	N
11	72	f	Yellow jacket	IV	F,C
12	56	f	Yellow jacket	IV	F,C
13	76	f	Yellow jacket	П	E, I
14	26	f	Yellow jacket	П	E,I
15	66	m	Yellow jacket	III	N,C
16	41	f	Yellow jacket	IV	F,C
17	35	f	Yellow jacket	П	F,C
18	34	m	Hornet	IV	F,C
19	69	f	Yellow jacket	III	F,I

or phentolamine [5]. In all published cases, no case of digital loss or other permanent damage was observed, irrespective of treatment. A retrospective cohort study on cases reported to six poison centers during six years found 365 epinephrine injections to the hand, of those, 213 were digital injections. All patients showed no significant systemic effects and complete resolution of symptoms [1]. Even in untreated patients, no necrosis or skin loss was described [5].

In our interview series, 89.5% of a cohort of 19 recently instructed patients were still carrying their autoinjector with them. 71.4% of our examined patients were capable to use the training device in a correct manner. In a study by De-Muth et al [6], only 37/63 food allergic-children were still carrying the prescribed autoinjector. In another study of 96 patients with insect venom allergy, less than 30% carried the autoinjector at all times and only 44% of patients were able to demonstrate proper autoinjector administration technique [7]. In a recently published study by Brown et al. [8], 15% of mothers without previous epinephrine autoinjector knowledge or experience could not "fire" training devices correctly despite a prior one-to-one demonstration of 30 minutes duration. Only 22% of mothers were able to perform all ten procedures according to predetermined criteria used to assess the mother's performance in using a randomly assigned epinephrine autoinjector device following a standardized teaching session by one clinician.

In another study by Topal *et al.* [9], 59/64 patients (14 children with food or venom allergy and 50 caregivers) obtained the autoinjector device after initial prescription. 42 (71%) still had the device at the time of the study. 54.6% of patients were not having an autoinjector because they felt that it was no longer necessary. There was a significant relation between adrenaline autoinjector competency and regular visits to the allergist. Autoinjector competency decreased as time elapsed from the last visit and the first instruction.

Arga and colleagues [10] demonstrated the importance of autoinjector design on unintentional injection injury. They compared a new and an originally designed epinephrine autoinjector with respect to correct use by 180 interns of the 2011-2012 training period of the medical school, randomly assigned into two groups. The number of participants correctly showing the use of epinephrine autoinjectors did not differ significantly between the two groups, but significantly fewer participants had unintentional injections in the new compared with the old epinephrine autoinjector group (p < 0.001). They concluded that the new epinephrine autoinjector injuries than the old one; however, it still did not fulfill the criteria of an ideal epinephrine autoinjector.

Overall, our observational study in accordance with previously published studies demonstrated that the compliance of carrying autoinjectors at all times and the ability of correct administration were poor and needed optimization.

In conclusion, a continuous, repetitive instruction of correct epinephrine autoinjector use by health care professionals is required.

CONFLICTS OF INTEREST

The authors confirm that this article content has no conflicts of interest.

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