An outbreak of plague occurred in the region of Oran, Algeria, from June to July 2003. Algeria had not reported this disease for >50 years. Eighteen bubonic cases were identified, and *Yersinia pestis* was isolated from 6 patients. Except for the index case-patient, all patients recovered. Targeted chemoprophylaxis, sanitation, and vector control played a crucial role in controlling the outbreak. Epidemiologic and biomolecular findings strongly suggested the existence of a local animal reservoir during this period, but its origin (resurgence or re-importation) could not be determined. This sudden and unexpected reemergence of plague, close to an important commercial seaport, is a textbook illustration of a public health event of international importance. It also demonstrates that the danger of plague reoccurrence is not limited to the currently indexed natural foci.

Plague is primarily a bacterial zoonosis affecting rodents. It is caused by *Yersinia pestis* and is transmitted from animal to animal by fleas. Humans usually become infected through the bite of an infected rodent flea. Bubonic plague, a severe infectious disease which, in the absence of appropriate antimicrobial drug therapy, can evolve to a rapidly fatal septicemia or pneumonia, can develop. A pneumonia form, which enables direct transmission to contacts, can be responsible for highly lethal outbreaks.

Currently, plague natural foci persist in Asia, the Americas, and Africa (where most human cases occur) (1). Plague foci have previously existed in the northern part of Africa but gradually disappeared in the last century, for unknown reasons. Libya is the only North African country that has experienced human cases in the past 40 years (2). In Algeria, archives report epidemics of plague as far back as the 14th century. These epidemics mainly affected ports, particularly that of Oran in 1556 and 1678 (3,000 deaths). In 1899, after an absence of nearly 100 years, plague reappeared in the port of Philippeville (now Skikda). Three large epidemics were subsequently reported in 1921 (185 cases), 1931 (76 cases), and 1944 (95 cases) as well as 158 sporadic cases. All but 2 cases occurred in ports (3,4). No natural focus of plague had ever been described in Algeria (5). We describe an outbreak of bubonic plague that occurred in 2003 in Algeria, where the last reported human case occurred in Oran in 1946 (6).

**Methods**

During June 9–18, 2003, several patients with signs of severe infection and painful inflammatory adenopathy were admitted to the University Hospital of Oran. All came from Kehailia (35°29′N, 0°32′E), a village of 1,300 inhabitants 25 km south of Oran. After eliminating all other possible differential diagnoses, clinicians suspected plague. The diagnosis was confirmed on June 18 by results of analysis of a bubo (lymph node) aspirate. A technical crisis committee was set up, and a case definition was adopted (Table). Any patient with a febrile syndrome and adenopathy who lived in the prefecture of Oran was hospitalized.

Clinical samples collected from patients (blood, bubo aspirate, cerebrospinal fluid) were sent to the Microbiology Department, University Hospital, Oran. Several of the initial cases were first diagnosed with the rapid diagnostic test (RDT) for plague developed by the Institut Pasteur (7); however, all samples were also examined with standard bacteriologic methods. Direct examination of smears was performed after Wayson and Gram staining. Blood samples were cultured in Castaneda medium for at least 10 days.
at 28°C and examined daily. Suspected samples were inoculated into brain heart infusion and peptone broth and streaked on blood agar and cefsulodin-irgasan-novobiocin (Merck, Rahway, NJ, USA) plates. All media were incubated at 28°C. Bacterial identification was conducted with API 20 E strips (Analytab Products, Syosset, NY, USA) or individual tests in tubes. The biovar was determined (8). Antimicrobial drug susceptibility testing (ampicillin, amoxicillin-clavulanic acid, cefazolin, cefotaxime, gentamicin, amikacin, sulfamethoxazole, doxycycline) was conducted according to the technique of the Clinical and Laboratory Standards Institute (www.clsi.org). The serodiagnosis was determined by the ELISA-F1 technique (9). Serum samples from 30 study participants who had not contracted the disease but lived in the same area as the patients were used to determine the positive threshold of the technique. A serum was regarded as negative if its optical density at 490 nm (OD_{490}) was lower than a threshold defined as the mean (M) OD_{490} value of normal sera + 3 standard deviations (SD): OD_{490} < M + 3 SD. Sera with OD higher than this threshold were regarded as weak when the ratio R = OD_{490}/(M + 3 SD) was <2 and positive if R was ≥2.

### Results

On June 9, 2003, a 19-year-old shepherd living in Kehailia was hospitalized with signs of septic shock (patient no. 2) (online Appendix Table, available from www.cdc.gov/EID/content/13/10/1459-appT.htm). He had been treated at home unsuccessfully with cephalosporins for inguinal adenopathy and fever during the previous 8 days. In the same village, 6 similar cases (nos. 3–8) occurred in the following days, until the diagnosis of plague was suspected. Five cases of bubonic plague (nos. 10, 11, 14, 15, and 17) subsequently occurred from June 21 to July 16 among persons living in villages around Kehailia.

On June 28, a farmer and his wife (patients 12 and 13) who resided in Ain Temouchent, 50 km west of Kehailia (Figure), were hospitalized in Oran for symptoms suggestive of plague. The patients reported that they had not left their farm during the weeks preceding their illness. On July 1, a child from Beni Saf, on the Mediterranean coast 100 km southwest of Kehailia (Figure), had clinical signs of bubonic plague and a positive RDT result (patient 16). Neither he, nor his parents, had gone to the area of Kehailia or Ain Temouchent during the previous days. The last case (patient 18) occurred on July 22. The patient, a hunter who lived in Oran, had walked in the forest of M’ila, 30 km northwest of Kehailia, a few days before onset of his clinical signs.

Altogether, 18 cases were identified June 4–July 22, 2003: 10 confirmed, 3 probable, and 5 suspected (or 12 confirmed, 2 probable, and 4 suspected, according to the new World Health Organization case definition [J]). Most of the patients lived in unsanitary conditions, in close contact with livestock, and in the vicinity of storage areas of grain and fodder. In Kehailia, all the case-patients resided in different dwellings located within a 200-m radius. None of them reported direct contact with rodents. Sixteen of the 18 patients had an inguinal bubo, indicative of a flea bite on the leg. A septicemic form of plague developed in patients 1 and 2. Patient 1 died very soon after hospital admission. Patient 2 was admitted with a severe fever and neurologic syndrome and fell into a deep coma, despite broad-spectrum antimicrobial drug treatment that included vancomycin, cefotaxime, and gentamicin. He recovered from the coma 48 hours after treatment with ciprofloxacin (500 mg 2×/d for 30 days) was completed (F. Razik et al., unpub. data). No case of secondary pulmonary dissemination was observed. Other plague patients were treated with either doxycycline for adults (200 mg/d for 10 days) or cotrimoxazole for children (40 mg/kg/d for 10 days). All recovered without sequelae.

On the whole, 60 bubo aspirates, 143 blood samples, 6 sputum samples, and 2 cerebrospinal fluid samples were analyzed. In 5 samples, smear stains suggested infection...
with *Y. pestis* (online Appendix Table). Among the 18 patients, 12 had a positive RDT result, but *Y. pestis* was isolated from only 6 patients: 5 from bubo aspirates and 1 from the blood culture of a patient whose bubo was too small to be punctured (patient 13). Results of ELISA-F1 serologic test conducted on the serum samples from 15 of the 18 patients were strongly positive 3 times and slightly positive 3 times (online Appendix Table).

**Discussion**

Epidemiologic investigation did not identify any other plague patients before patient 1. It is unlikely that other cases occurred and remained undetected during this period since plague, even in its bubonic form, is a severe infection with high fatality rates.

For the first time, the RDT was used in an epidemic situation outside of Madagascar, where it was developed. The case definition had to take into account this particularity. The bacteriologic diagnosis is a long procedure (at least 4 days) and, in this epidemic context, RDT contributed to the effectiveness of the response. Of the 44 RDTs that were conducted, 12 had positive results; by contrast, culture was positive only for 6. Among the 15 patients for whom a serologic test was conducted (online Appendix Table), a specific antibody response developed only in 6. This absence of specific antibodies can be explained by the fact that serum specimens were taken before the appearance of anti-F1 immunoglobulin G, or by a rapid administration of antimicrobial drugs, which stopped development of an immune response. The 3 clearly seropositive patients were those from whom a positive culture was obtained.

The outbreak occurred in a poor rural settlement, with inadequate sanitation. The residents observed an increase in the population of commensal rodents, which is often associated with the harvesting period, but no unusual rodent mortality was noted during the weeks preceding the outbreak. The appearance during the same week of 2 new cases in Ain Temouchent (50 km west of Kehailia) and then 1 case in Beni Saf (100 km southwest of Kehailia) could not be explained. Nonetheless, the fact that the *Y. pestis* strains isolated in Kehailia and Ain Temouchent had identical pulsortypes (V. Chenal-Francisque et al., unpub. data) argues for a single focus and not for independent foci that emerged simultaneously.

A crisis committee designed and supervised a control strategy based on standardized case management, prophylactic treatment and follow-up of contacts sharing the same dwelling as plague patients, and vector control. Environmental sanitation measures in Kehailia contributed to reduction in the occurrence of new cases in this village. Intra- and peridomestic spraying with permethrin was conducted. Deltamethrin was dusted on the tracks and around the burrows of rodents located in a radius of 10 km around the dwelling of the patients. Uncontrolled killing of rats was prohibited.

No natural focus of plague had ever been described in Algeria. Past cases were always regarded as imported through the ports. The reappearance of human cases in this area can be explained in 2 ways: a recent importation of infected animals or a sudden manifestation of a natural focus that had remained silent for decades. It is noteworthy that Kehailia, the epicenter of the outbreak, is in the vicinity of flour mills built 4 years before the outbreak. These mills are supplied regularly with cereals by trucks arriving from the port of Oran. A part of this traffic was still run by railway a year before the outbreak, and a marshalling yard was installed a few kilometers from Kehailia. In 1919, this mode of importation was responsible for the plague outbreak that
occurred 75 km south of the port of Skikda (10). The hypothesis of recent importation of the plague bacillus in Kehailia is therefore tempting but is tempered by the fact that 1) the grain is primarily imported from Europe, which is not affected by plague, and from North America where natural foci exist but have very limited areas of overlap with those regions where cereal grains are grown, 2) no higher mortality rate in the murine population of the port was noted, 3) no human cases occurred in this sector of the city, and 4) a 3IS–restriction fragment length polymorphism (11) analysis grouped these strains in a cluster clearly distinct from the strains isolated from Africa and America (V. Chenal-Francisque et al., unpub. data).

The geographic concentration of the cases in 2 foci, both contiguous in the mountainous area of Tessala, suggested the existence of a natural focus in this area. Moreover, Meriones are present in Tessala, and these rodents are a well-known potential reservoir of Y. pestis (12). The outbreak occurred at harvest time, and it is possible that the abrupt reduction in the source of food pushed the wild rodents to approach houses in which grain was stored.

The current challenge in terms of public health is to determine if this animal reservoir has disappeared or if it is well established in the ecosystem. The capture of 3 se-ropositive small mammals (2 Mus musculus and 1 Aleterix algerius) in July 2004 (J.L. Soares et al., unpub. data) and the identification of several Y. pestis infected fleas in the same area (13) favor the second option.

Beyond the local problem, the proximity of a possible natural reservoir of plague to Oran, a large international commercial port, raises the possibility of the risk for an urban outbreak. At the time of the investigation, the sanitation in the city and port were poor and rodents proliferated. These urban rodents could come in contact with infected rodents from rural areas in the uncontrolled dumps at the periphery or through a dry riverbed that penetrates as far as the city center. Because of Oran’s population density and the commercial activities of its seaport, a plague outbreak would have international implications.

This outbreak is a textbook illustration of the unexpected and sudden reemergence of an infectious disease epidemic that is potentially highly lethal. It also demonstrates that the danger of a plague outbreak is not limited to the currently indexed natural foci.

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“So many scientists think that once they figure it out, that’s all they have to do, and writing it up is just a chore. I never saw it that way; part of the art of any kind of total scholarship is to say it well.”

—Stephen Jay Gould
### Appendix Table. Characteristics, clinical manifestations, and laboratory results of plague patients in Algeria, June–July 2003*

| Patient no. | Sex | Age, y | Place of residence | Occ. | Date of admission | Clinical manifestations | Evolution | Direct examination | Culture | RDT | Sero 1 (bleeding date) | Sero 2 | Remark | Class. |
|-------------|-----|--------|---------------------|------|-------------------|------------------------|-----------|-------------------|----------|-----|-----------------------|--------|---------|--------|
| 1           | M   | 11     | Kehailia            | Pupil| Jun 4             | Septicemia             | Died       | –                 | –        | –   | –                     | –      | Cousin of patient no. 2 | S       |
| 2           | M   | 19     | Kehailia            | Shepherd| Jun 9          | Septicemia            | Recovered   | Neg               | Neg      | Pos | Neg (Jul 12)          | –      | Cousin of patient no. 1 | C       |
| 3           | M   | 70     | Kehailia            | Farmer| Jun 13           | Inguinal bubo          | Recovered   | Neg               | Neg      | Pos | Weak (Jul 7)          | –      | Head of the village   | C       |
| 4           | F   | 27     | Kehailia            | None  | Jun 14           | Inguinal bubo          | Recovered   | Pos               | Pos      | Pos | –                     | –      | –                   | C       |
| 5           | M   | 26     | Kehailia            | Shepherd| Jun 15         | Inguinal bubo          | Recovered   | Neg               | Neg      | Pos | –                     | –      | –                   | C       |
| 6           | M   | 20     | Kehailia            | Trader| Jun 16           | Inguinal bubo          | Recovered   | –                 | –        | –   | –                     | –      | –                   | S       |
| 7           | M   | 20     | Kehailia            | None  | Jun 17           | Inguinal bubo          | Recovered   | –                 | –        | –   | –                     | –      | –                   | S       |
| 8           | F   | 54     | Kehailia            | None  | Jun 17           | Inguinal bubo          | Recovered   | –                 | –        | –   | –                     | –      | Stay in Kehailia       | S       |
| 9           | F   | 25     | Hai Oussama         | None  | Jun 19           | Inguinal bubo          | Recovered   | Neg               | Pos      | Pos | Neg (Jul 7)           | –      | 3 km from Kehailia     | C       |
| 10          | M   | 27     | Karma Es-Senia     | Farmer| Jun 21           | Inguinal bubo          | Recovered   | Pos               | Neg      | Neg | Neg (Jul 7)           | –      | 2 km from Kehailia     | S       |
| 11          | M   | 55     | Hama Ali            | Mason | Jun 27           | Inguinal bubo          | Recovered   | Pos               | Pos      | Pos | Pos (Jul 8)           | Pos    | Husband of 13          | C       |
| 12          | M   | 58     | Ain Temouchent      | Shepherd| Jun 28         | Inguinal bubo          | Recovered   | Pos               | Pos      | Pos | Pos (Jul 21)          | Pos    | Wife of 12             | C       |
| 13          | F   | 55     | Ain Temouchent      | None  | Jun 28           | Inguinal bubo          | Recovered   | –                 | Pos      | –   | Weak (Jul 21)         | –      | 10 km from Kehailia    | C       |
| 14          | F   | 2      | Zeghloul            | None  | Jun 30           | Inguinal bubo          | Recovered   | Neg               | Neg      | Pos | –                     | –      | 4 km from Kehailia     | C       |
| 15          | F   | 3      | Hamoul              | None  | Jun 30           | Inguinal bubo          | Recovered   | Pos               | Pos      | Pos | Pos (Jul 8)           | Pos    | 4 km from Kehailia     | C       |
| 16          | M   | 7      | Beni Saf            | None  | Jul 1            | Inguinal bubo          | Recovered   | Neg               | Neg      | Pos | Neg (Jul 7)           | –      | 25 km from Kehailia    | P       |
| 17          | F   | 24     | El Karma            | Civil servant | Jul 16        | Axillary bubo          | Recovered   | Neg               | Neg      | Pos | Weak (Jul 20)         | –      | Stay in the house of 8 | P       |
| 18          | M   | 20     | Oran                | Hunter| Jul 22           | Inguinal bubo          | Recovered   | Neg               | Neg      | Pos | –                     | –      | –                   | P       |

*Occ., occupation; RDT, rapid diagnostic test; Class., classification; –, not done; Neg, negative; Pos, positive; C, confirmed case; S, suspected case; P, presumptive case.