

ROENTOROLL 25

Application

Specially designed for table processors and processors with small tank volumes, such as the Agfa Gevamat 60, Agfa Curix 60, Curix Compact, Scopix Compact, Dürr AC 245 L, Medicine 260, Fuji RG II, FPM 100 A, Kodak ME 10.

ROENTOROLL 25 is suitable for all areas of X-ray diagnostics, including dental film processing and X-ray film processing for non-destructive testing (NDT).

Characteristics

All the standard X-ray films are developed with a good speed yield and high contrast. **ROENTOROLL 25** features great stability, and so ensures consistent sensitometric results over lengthy periods, even with processors run at a low film throughput.

ROENTOROLL 25 does not need to be mixed with starter.

The concentrates will keep for at least 24 months at storage temperatures between 5°C and 30°C.

Pack sizes

ROENTOROLL 25 is a two-part developer concentrate, and is supplied in a pack for 12 x 2.5 litres of ready-to-use developer. Due to the good ratio of concentrate to ready-to-use solution, the weight ratio for a pack unit is very good.

Developer temperature

The developing temperature depends on the X-ray film used and on the immersion time in the developer, which varies according to processor type.

The standard figures are:

31 - 33°C	at 2 min total processing time
27 - 30°C	at longer total processing times

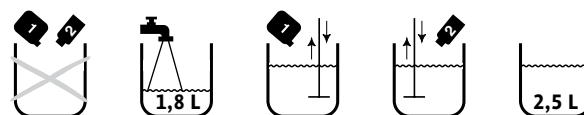
Replenishment

ROENTOROLL 25 should be replenished at 40 ml to 65 ml per 35 x 35 cm sheet. The replenishment rate required for a processor depends on the film throughput. If throughput is low and there are long intervals between processing, the higher replenishment rate must be used. If throughput is high the lower replenishment rate can be used.

Mixing instructions

Never mix Part 1 and 2 concentrates!

Use 1.8 litres of water. Parts 1 and 2 are stirred in, and produce 2,5 litres of ready-to-use developer. **No starter is needed.**



Certification

Tetenal operates a fully certified quality management system to DIN EN ISO9001.

This product complies with the medical devices directive 93/42/EEC concerning medical products (CE-marking).