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SpeedUpMyPC 3.5 Cracked Serial-maz Serial Key Download-Keygen Code. Download From Our Website : Materizer.me Visit us : <http://1>. Field of the Invention The present invention relates to a novel buffer material. More specifically, the present invention relates to a novel buffer material for the preservation of biological specimens. 2. Description of the Related Art In order to preserve a biological specimen of a specific origin, the specimen is generally treated by incorporating the specimen in a preservative medium or a buffer material. Furthermore, in order to stabilize a specific ionic compound contained in the biological specimen, a buffer material may be added to the preservative medium. The buffer material functions to the preservation of a biological specimen by neutralizing an acidic substance (for example, a protease) contained in the specimen to prevent the degradative enzyme from breaking down the specimen. Conventionally, an inorganic buffer material (for example, sodium carbonate) has been used as the buffer material. The inorganic buffer material, however, has a deficiency in that the buffer capacity thereof is not high. In addition, the inorganic buffer material is not appropriate for preserving blood or tissues, which is composed of approximately 65% of water. However, the inorganic buffer material has an advantage in that it does not dissolve in an organic solvent, and therefore, the biological specimen does not contact with the organic solvent. On the other hand, the inorganic buffer material has a disadvantage in that it is readily soluble in water. Accordingly, the inorganic buffer material is not suited for a preservative medium for preserving biological specimens in a blood or tissue, which is mainly composed of water. Furthermore, most biological specimens are degraded by a protease contained therein. Accordingly, in order to effectively use the biological specimen, the protease must be inactivated. In view of this fact, a variety of protease inhibitors have been developed and used. In general, a protease inhibitor has a structure including a substrate-recognizing group (substrate binding site) capable of inhibiting a protease and a binding site for binding the protease. The protease inhibitors have a plurality of functional groups capable of binding to the protease, respectively, and thus, the inhibitors can bind to a plurality of proteases. However, each protease inhibitor has a tendency to bind to only one protease, and therefore, it is difficult to inhibit more than one protease. As a result,

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