

the stand is in the assembled configuration, the back support defining first and second support slots and comprising a foldably positionable first arm and a foldably positionable second arm, each of said first and second arms comprising a flap tab and a support tab, wherein the first and second arms are substantially parallel with said upper surface of said base when the stand is in the folded configuration, and are substantially perpendicular to said upper surface of said base when the stand is in the assembled configuration, and further wherein the first and second arms are configured to be positioned on the upper surface of said base when the stand is in the assembled configuration; wherein the flap tab of the first and second arms is configured to reversibly fit into the first and second flap slot, respectively, of the upper support front flap when said stand is in said assembled configuration, and further wherein the support tab of the first and second arms is configured to reversibly fit into the first and second support slot, respectively, when said stand is in said assembled configuration.

10. The stand of claim 9, wherein said stand comprises cardboard.

11. The stand of claim 9, wherein the upper platform defines a back support slot and said back support comprises a back support tab, the back support tab configured to fit into the back support slot when said stand is in said assembled configuration.

12. The stand of claim 9, wherein said base defines a cavity.

13. The stand of claim 9, wherein said upper platform comprises a notch sized to receive at least a portion of an umbilical cord.

Description

BACKGROUND OF THE INVENTION

The present invention relates to procurement of cord blood and placental stem cells and, more specifically, to devices and systems, and methods for manipulating and holding a placenta during procurement of cord blood and placental stem cells.

Stem cells are master cells found in all multicellular organisms. These special cells are important to the human body, for example, because they are capable of differentiating into a multitude of different specialized cell types, and dividing to maintain a supply of stem cells. In humans there are two main types of stem cells: embryonic stem cells and adult stem cells. In a developing embryo stem cells differentiate into all types of cells, thereby creating specialized tissues, organs, and systems. In an adult human, stem cells are involved in the normal turnover of organs such as blood and skin.

Hematopoietic stem cells, for example, are used to treat blood and immune system diseases because they can differentiate into red blood cells, white blood cells, and platelets. However, some stem cell transplants have been performed for patients with genetic or metabolic diseases. Indeed, to date more than 80 different diseases have been treated using stem cell transplants. According to the National Cord Blood Program, there were over 15,000 through the end of 2009. The National Marrow Donor Program estimates that there will be 10,000 cord blood transplants per year by 2015, up from 2,000 per year in 2006.

In addition to known treatments involving stem cells, research continues into the promise of many potential future applications. Indeed, the ability of stem cells to differentiate into other types of cells holds significant promise for treating some of the world's most common diseases including heart disease, diabetes, stroke, hearing loss, blood disorders, Parkinson's disease, and Alzheimer's disease, just to name a few.

Umbilical cord blood--blood which remains in the placenta and umbilical cord after childbirth--is one of the most common sources of stem cells. Since cord blood is collected from the placenta, which is normally discarded, the collection process is safe for both the mother and the newborn. Cord blood is obtained by syringing out the placenta through the umbilical cord shortly after childbirth, after the cord has been detached from the newborn. The retrieved blood can then be frozen and stored indefinitely.

FIG. 8, for example, shows a collection kit 600 (schematic shown in FIG. 4) that has been placed inside the pocket or cavity 560 formed in the back of base 110. The edges of the pocket are shown around the collection kit 600.

Accordingly, to assemble a stem cell collection stand from a first, folded configuration to a second, assembled configuration, the following steps are performed. First, a folded stand 100 is provided, as shown in FIG. 7. Next, the folded components are removed from the case 200, as shown in FIGS. 8 and 9. Next, the upper portion 130 is pulled upward, and rear support 340 (which is connected at its base to the base 110) is pulled upward, hinging at fold 510 so that backing 130 is vertical between the base 110 and upper portion 130, as shown in FIG. 10, with tab 400 of rear support 340 fitting into slot 410 of upper portion 130. Arms 310 and 320 are pulled outward from the flat configuration in FIG. 9 to the extended configuration (see arm 320 in FIG. 10). Front flap 390 of upper portion 130 is folded underneath so that the flap faces the interior of the vertical portion of the device, as shown in FIG. 10. Each of arms 310 and 320 are folded backward toward backing 130, such that tabs 350 of each arm fit into slots 370 of the folded front flap 390 of upper portion 130, and tabs 360 of each arm fit into slots 380 of rear support wall 340, as shown in FIG. 10. Device 100 is now in the second, assembled configuration shown in FIG. 1.

In addition to facilitating the ingress and egress of stem cell procurement fluids into and out of a placenta, maximizing efficient stem cell collection also often requires the secure transport of the placenta from one location to another. According to one embodiment, the kit can also include a shipping container 700, as shown in FIG. 6. The shipping container can be utilized to ship one or more components of the device and/or collection quantities (stem cells, placenta, cord blood, etc.) to a facility for storage or processing.

Although the present invention has been described in connection with a preferred embodiment, it should be understood that modifications, alterations, and additions can be made to the invention without departing from the scope of the invention as defined by the claims.

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