Diagnostic Algorithm in Hirschsprung’s Disease: Focus on Immunohistochemistry Markers

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Abstract. Background/Aim: Hirschsprung disease (HD) is caused by the congenital absence of ganglion cells in the distal bowel (aganglionosis). Rectal biopsy is considered important for its diagnosis. The aim of this study was to apply immunohistochemical staining using a minimal set of antibodies and develop an algorithm that will assist in the diagnosis of HD. Patients and Methods: Rectal or colonic biopsies were performed in patients with HD (n=26) or patients treated for other bowel diseases (n=34). Immunohistochemical staining was performed for MAP1b, peripherin, S-100, calretinin, NSE, bcl-2 and CD56 proteins. Results: Staining for CD56, S-100, peripherin and calretinin facilitated the identification of ganglion cells. The use of CD56 and S-100 antibodies together resulted in the highest rate of ganglion cell staining intensity (94%). Conclusion: We propose a practical diagnostic algorithm with the application of CD56 and S-100 antibodies that can be used in clinical practice in children suspected of Hirschsprung’s disease.

Hirschsprung disease (HD) is a congenital, genetically-based functional obstruction due to the congenital absence of ganglion cells in the distal bowel (aganglionosis) (1, 2). The absence of ganglion cells along with an analysis of hypertrophy and hyperplasia of nerves in the nerve plexus of submucosa and muscularis mucosae in rectal biopsy specimens (RB) are regarded as potential hallmarks for its diagnosis (3). Hematoxylin and eosin (HE) staining alone in the assessment of the presence of ganglion cells (GC) is still challenging, especially in newborns with total colonic aganglionosis (TCA) or intestinal neuronal dysplasia (IND) (4). Diagnosis of the disease is mainly based on the identification of the lack of ganglion cells in the pathology sections of the colon and often requires an additional immunohistochemical (IHC) staining, which is very difficult and time consuming and also needs several serially cut sections (5, 6).

This study was performed in order to ascertain the role of IHC staining with a selected set of antibodies in diagnosing HD and develop a diagnosis algorithm based on the selection of a minimal set of antibodies sufficient for its diagnosis.

Patients and Methods

Patients. A total of 60 children were enrolled in the study, including 26 with clinical symptoms of aganglionosis (HD, Group 1) and 34 with other diseases, such as meconium ileus, necrotizing enterocolitis, chronic constipation or sigmoid volvulus (non-HD, Group 2). All patients with aganglionosis (Group 1) and 24/34 with non-HD (Group 2) were male. Median age was the same in both groups: 9 months (range=6-30 months) and 73% and 76% were under 1 year, respectively. In all cases, rectal or colonic biopsies were performed and the specimens were archived in formalin-fixed paraffin-embedded tissue sections, followed by HE and IHC staining.

Immunohistochemistry (IHC). The IHC staining was performed using a set of antibodies against: MAP1b neuronal marker (Abcam, Cambridge, UK), peripherin (Novocastra, Newcastle, UK), S-100
(DAKO, Glostrup, Denmark), calretinin (DAKO), NSE (DAKO), bcl-2 (DAKO) and CD56 (DAKO). To determine the appropriate antibody dilution to eliminate false-positive results, as well as to reduce background reactions, a series of control reactions were performed before adequate immunohistochemical staining. Positive and negative control reactions were performed in each case. The independent assessment was performed by two experienced pathologists.

In order to identify the best set of antibodies for GC identification in IHC staining, the following four criteria were used: 1. possibility of GC distinction from other neural components; 2. lack of artifacts; 3. good intensity of GC staining; 4. the highest fulfillment rate comparing results of different antibodies. These tests were performed on non-HD samples (Group 2).

To assess the possibility of GC distinction from other neural components (first criterion), a GC distinction index was created: GC distinction=$(0\times n_a+1/2\times n_m+n_e)/(n_a+n_m+n_e)\times100\%$, where: $n_a$ – number of samples where GC were absent; $n_m$ – number of samples where GC were moderately easy to detect; $n_e$ – number of samples where GC were easy to detect. Assessment of GC staining intensity was semiquantitative using 4 groups of results: no staining or artifacts, weak staining of GC, GC well visible and evidently visible. To assess good intensity of GC staining (third criterion), a GC staining intensity index was created: GC staining intensity=$[(0\times n_0+1/3\times n_1+2/3\times n_2+n_3)/(n_0+n_1+n_2+n_3)]\times100\%$, where: $n_0$ – number of samples with no staining or artifacts; $n_1$ – number of samples where GC were weakly stained; $n_2$ – number of samples where GC were moderately stained; $n_3$ – number of samples where GC were evidently stained. The study was approved by the Institutional Review Board, (Department of General and Oncological Surgery for Children and Adolescents, KB 167/2012).

Statistical analysis. Categorical variables were compared with the chi-square or Fisher exact test, and non-categorical variables were compared with the Mann-Whitney U-test.

Results

Four antibodies clearly facilitated the identification of ganglion cells (first criterion) in the IHC studies: CD56, S-100, peripherin and calretinin with a more than 60% index of GC distinction (Figure 1A).

Analysis of the rate of artifacts (second criterion) revealed that anti-S-100 and anti-bcl-2 antibodies were the most efficient (0% of negative staining). A relatively high rate of negative staining was observed for MAP1B (31%) and calretinin (18%), while the best intensity of GC staining (third criterion) was found for CD56 (91%) and peripherin (83%) (Figure 1B). The overall sensitivity, specificity, positive, and negative predictive value of antibody staining is shown in Table I.

The fourth criterion was assessed using cluster analysis and the largest incompatibility rate (Table II). Two groups of results were defined: the first cluster included MAP1B, S-100 and bcl-2, while the second one included peripherin, CD56, NSE and calretinin. As only four antibodies clearly facilitated the identification of ganglion cells (Figure 1A), available sets of antibodies included: CD56 and S-100, peripherin and S-100, and calretinin and S-100 with summary marker expression indexes: 94%, 88% and 74%, respectively. The identification of ganglion cells was the most efficient for the set of CD56 and S-100, in cases without artifact staining.

In order to verify the value of IHC in patients with aganglionosis, an expanded set of IHC studies was performed on biopsy specimens obtained from patients with Hirschsprung’s disease (Figure 2). The staining with CD56, S-100, peripherin and calretinin demonstrated the presence of GC in 3/26 patients initially qualified as aganglionosis in HE examination. In two patients, the morphology of GC was truly dysplastic, and the third patient was further diagnosed and treated for food allergy. All 23/26 HD cases after surgical treatment were uncomplicated, except for more frequent bowel movements and perianal excoriations for a period of 1-2 months in two patients.
Discussion

Nowadays, surgical treatment of Hirschsprung disease is performed at an earlier age and more often in a one-stage manner and therefore, requires very precise preoperative diagnosis (7). HE staining of rectal biopsy specimens is a gold standard for HD and differential diagnosis (2, 8, 9). Proper assessment of GC presence requires primarily adequate tissue sampling and careful tissue preparation. There are numerous reports on the use of various antigens in IHC studies for better diagnosis of aganglionosis. An ideal immunohistochemical marker would facilitate easy detection of GC through good, intensive staining, as well as easy distinction of GC from other neural components, and a low artifact rate (10).

We showed that selection of a pair of IHC assay reagents facilitates the diagnostic examinations of patients tested for HD. In the meantime, high sensitivity, specificity and efficacy of peripherin and S-100 staining was confirmed in a diagnostic protocol for HD. Ganglion cells were diagnosed in 93% of samples excluded in the primary examination (11). According to our algorithm, rectal biopsies with ambiguous histopathological presentation that does not allow for confirmation or exclusion of HD were subject for internal verification as assessed by another experienced pathologist, and final diagnosis was pursued by means of the proposed set of IHC assays.

A biopsy for the ganglion cells, and consequently, for neural hypertrophy is regarded to be the basic modality for the diagnosis of HD. S-100 staining was shown to be diagnostic in rectal suction biopsies in the samples that were devoid of ganglion cells. Staining for S-100 coupled with calretinin and PGP9.5 immunostaining performed on suction rectal biopsies was shown to be sensitive and specific for diagnosing HD (12). However, S-100 was found to be positive in ovarian cancer (13), metastases to the small intestine (14) and fibrosarcoma cells (15). The other marker, CD56, was used in evaluating a “transition zone” proximally to the aganglionic segment, with the use of serial colonic wall biopsies and obtained very good results (16). CD56 is known to be useful for definitive histopathological diagnosis of isolated hypoganglionosis. In the aganglionic segment in HD patients, CD56-positive extrinsic nerve fibers and bundles replaced typical ganglion cells that are immunoreactive for Hu C/D (17). Other recommended combinations of staining included AChE and calretinin (6) or peripherin and calretinin (18).

The practical value of the study is related to clinical verification: in three cases classified as HD, the use of the examined panel of immunohistochemical reagents allowed for clear visualization of ganglion cells, and therefore for verification of the primary diagnosis of HD. The histopathological diagnostic algorithm presented in this study provides a tool for improving the sensitivity and specificity of assays used in the analysis of specimens and eventually for a reduction in the number of patients unnecessarily subjected to surgical treatment.

In conclusion, we created a practical diagnostic algorithm with the application of CD56 and S-100 markers that can be

| % | MAP1B | Peripherin | S100 | Calretinin | NSE | Bcl-2 | CD56 |
|---|---|---|---|---|---|---|---|
| Sensitivity | 26.5 | 91.2 | 97 | 76.4 | 97 | 97 | 91.2 |
| Specificity | 95.6 | 95.6 | 100 | 100 | 95.6 | 100 | 100 |
| PPV | 90 | 96.8 | 100 | 100 | 97 | 100 | 100 |
| NPV | 46.8 | 88 | 95.8 | 74 | 95.6 | 95.8 | 88.4 |

PPV: Positive predictive value; NPV: Negative predictive value.
used in IHC in clinical practice for the diagnosis of HD. Nevertheless, additional studies are needed to confirm the value of calretinin and peripherin in patients diagnosed for aganglionosis.

**Conflicts of Interest**

The Authors have no conflicts of interest to disclose in regard to this study.

**Authors’ Contributions**

PG, AM: concept/design, data analysis/interpretation; LS, MB: data collection, data analysis/interpretation; PG, AM, JS: writing, critical revision and approval of the article.

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